

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: February 19, 2004, 21:05:58 ; Search time 1571 Seconds
(without alignments)
324.885 Million cell updates/sec

Title: US-10-085-108-21_COPY_711_731

Perfect score: 21

Sequence: 1 AAGCTGATTGATCACCAGGG 21

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 22781392 seqs, 12152238056 residues

Total number of hits satisfying chosen parameters: 45562784

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:

1: em_estba:*
2: em_esthum:*
3: em_estin:*
4: em_estmu:*
5: em_estov:*
6: em_estpl:*
7: em_estro:*
8: em_hic:*
9: gb_est1:*
10: gb_est2:*
11: gb_hic:*
12: gb_est3:*
13: gb_est4:*
14: gb_est5:*
15: em_estfun:*
16: em_estom:*
17: em_gss_hum:*
18: em_gss_inv:*
19: em_gss_pin:*
20: em_gss_vrt:*
21: em_gss_fun:*
22: em_gss_mam:*
23: em_gss_mus:*
24: em_gss_pro:*
25: em_gss_rod:*
26: em_gss_phg:*
27: em_gss_vrl:*
28: gb_gss1:*
29: gb_gss2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	18.4	87.6	686	29	CC270269 CH261-1N2
2	18	85.7	809	29	BZ721194 PUCFH73TD
C 3	17.8	84.8	307	9	AA894450 oe79f07.s
C 4	17.8	84.8	339	9	AI572687 tr78b10.x

5	17.8	84.8	380	28	BH323391	BH323391 CH230-53J
C 6	17.8	84.8	545	14	CA941354	CA941354 ir34b01.x
C 7	17.8	84.8	592	9	AI816423	AI816423 au47g07.x
C 8	17.8	84.8	671	29	AG121419	AG121419 Pan ttogl
C 9	17.8	84.8	708	29	AG186976	AG186976 Pan ttogl
C 10	17.8	84.8	913	13	HU245702	HU245702 603783316
C 11	17.8	84.8	963	10	BG119393	BG119393 602349076
C 12	17.8	84.8	1201	9	AL523235	AL523235 AL523235
C 13	17.4	82.9	565	28	AZ072191	AZ072191 RPCI-23-3
C 14	17.4	82.9	914	10	BE729033	BE729033 601561212
C 15	17	81.0	543	9	AI818091	AI818091 wk27a09.x
C 16	17	81.0	733	14	CB983939	CB983939 AGENCOURT
C 17	17	81.0	763	14	CD101012	CD101012 AGENCOURT
C 18	16.8	80.0	321	9	AA967543	AA967543 ua07e06.r
C 19	16.8	80.0	341	9	AA118769	AA118769 mp61e01.r
C 20	16.8	80.0	348	29	AG236818	AG236818 Lotus jap
C 21	16.8	80.0	359	10	BE861602	BE861602 UI-M-AH1-
C 22	16.8	80.0	364	28	BH056156	BH056156 RPCI-24-3
C 23	16.8	80.0	369	9	AW229797	AW229797 uo61h04.y
C 24	16.8	80.0	409	9	AA636853	AA636853 vrl9h04.r
C 25	16.8	80.0	420	10	BE181117	BE181117 CM3-HT062
C 26	16.8	80.0	457	9	AI225508	AI225508 ue88d11.y
C 27	16.8	80.0	457	10	BF322594	BF322594 maa39b08.
C 28	16.8	80.0	497	10	BG276054	BG276054 uu89e08.y
C 29	16.8	80.0	509	9	AW230744	AW230744 uo67d02.y
C 30	16.8	80.0	512	13	BU583368	BU583368 mai04c04.
C 31	16.8	80.0	522	10	BG078216	BG078216 H3025A02-
C 32	16.8	80.0	523	13	BQ356299	BQ356299 CM3-HT011
C 33	16.8	80.0	526	13	BQ287718	BQ287718 ii68f07.y
C 34	16.8	80.0	543	9	AA798972	AA798972 vv95e04.r
C 35	16.8	80.0	544	4	EX517147	EX517147 RZPD Mus
C 36	16.8	80.0	554	9	AA543306	AA543306 vk36e11.r
C 37	16.8	80.0	556	9	AA543338	AA543338 vk36h11.r
C 38	16.8	80.0	563	14	W19670	W19670 zb36a03.r1
C 39	16.8	80.0	581	14	CB271143	CB271143 mai55g11.
C 40	16.8	80.0	583	18	AQ047644	AQ047644 CLM-289-u
C 41	16.8	80.0	587	13	BQ444442	BQ444442 UI-M-ER0-
C 42	16.8	80.0	593	13	BQ265035	BQ265035 NISC_ff03
C 43	16.8	80.0	600	12	BI987638	BI987638 3204-05 M
C 44	16.8	80.0	604	9	AA544735	AA544735 vk35d11.r
C 45	16.8	80.0	640	29	BZ421605	BZ421605 hz32b04.b

ALIGNMENTS

RESULT 1
CC270269
LOCUS CH261-1N21 Sp6.1 CH261 Gallus gallus genomic clone CH261-1N21, 686 bp DNA linear GSS 13-MAY-2003
DEFINITION Genomic survey sequence.
ACCESSION CC270269
VERSION CC270269.1 GI:30623166
KEYWORDS GSS.
SOURCE Gallus gallus (chicken)
ORGANISM Gallus gallus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Archosauria; Aves; Neognathae; Galliformes; Phasianidae;
Phasianinae; Gallus.
REFERENCE 1 (Bases 1 to 686)
AUTHORS Kremizki,C., Higginbotham,J., Wylie,K., Carter,J., McPherson,J., Warren,W., Graves,T., Mardis,E. and Wilson,R.
TITLE Gallus gallus BAC End Reads
JOURNAL Unpublished
COMMENT Contact: Richard K. Wilson
Genome Sequencing Center
Washington University School of Medicine
Email: submisson@wustl.edu
Insert Length: 182000 Std Error: 0.00
Seq primer: Sp6 ATTAGGTGACACTATAG
Class: BAC ends
High quality sequence start: 68
High quality sequence stop: 498.

```

FEATURES
source
  Location/Qualifiers
  1. .686
    /organism="Gallus gallus"
    /mol_type="genomic DNA"
    /strain="Red Jungle Fowl"
    /db_xref="taxon:3031"
    /clone="CH261-1N21"
    /sex="Female"
    /cell_line="UCD001, inbred 256"
    /clone_lib="CH261"
    /note="Vector: pTARBAC2.1; Site 1: EcoRI; Site 2: EcoRI;
    CH261 Female Chicken Library - For library and clone
    ordering information: http://www.chori.org/bacpac"
  BASE COUNT    196 a    147 c    117 g    226 t
  ORIGIN

  Query Match      87.6%; Score 18.4; DB 29; Length 686;
  Best Local Similarity 95.0%; Pred. No. 3.4e+02;
  Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

  QY  2 AGCTGATTGATGACACAGG 21
      |||||
  Db   524 AGCTGATTGAAGACACAGG 543

  RESULT 2
  BZ7211194
  LOCUS
  DEFINITION
    PUCFH73TD ZM 0.6 1.0 KB Zea mays genomic clone ZMMBta133M02,
    genomic survey sequence.
  ACCESSION
    BZ7211194
  VERSION
    BZ7211194.1 GI:28513208
  KEYWORDS
    GSS.
  SOURCE
    Zea mays
  ORGANISM
    Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
    Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACAD
    clade; Panicoideae; Andropogoneae; Zea.
  1 (bases 1 to 809)
  Whitelaw,C.A., Quackenbush,J., Van Aken,S., Utterback,T., Resnick
  A., Fraser,C.M., Yuan,Y., San Miguel,P., Ma,J. and Bennetzen,J.
  Maize Genomics Consortium
  Unpublished
  Contact: Cathy Whitelaw
  TIGR
    9712 Medical Center Drive, Rockville, MD 20850, USA
    Tel: 301-838-5843
    Fax: 301-838-0208
    Email: whitelaw@tigr.org
  Seq primer: TF
  Class: sheared ends.
  Location/Qualifiers
  1. .809
    /organism="Zea mays"
    /mol_type="genomic DNA"
    /strain="B73"
    /db_xref="taxon:4577"
    /clone="ZMMBta133M02"
    /note="Vector: PCR4-TOPO; Site 1: EcoRI; 0.6-1.0 kb high
    Cot selected genomic DNA library"
  BASE COUNT    231 a    173 c    172 g    233 t
  ORIGIN

  Query Match      85.7%; Score 18; DB 29; Length 809;
  Best Local Similarity 100.0%; Pred. No. 5.8e+02;
  Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

  QY  4 CCTGATTGATGACACAGG 21
      |||||
  Db   773 CCTGATTGATGACACAGG 790

```

```

RESULT 3
AA894450/c
LOCUS
DEFINITION
  oa79f07.s1 NCI_CGAP_Lu5 Homo sapiens cDNA clone IMAGE:1417861 3',
  mRNA sequence.
ACCESSION
  AA894450
  AA894450.1 GI:3030851
  EST.
  Homo sapiens (human)
  SOURCE
    Homo sapiens
  ORGANISM
    Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
    Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
  REFERENCE
    1 (bases 1 to 307)
    NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
    National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
    Tumor Gene Index
  JOURNAL
    Unpublished
  COMMENT
    Contact: Robert Strausberg, Ph.D.
    Email: cgaps-f@mail.nih.gov
    Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
    Emmert-Buck, M.D., Ph.D.
    cDNA Library Preparation: M. Bento Soares, Ph.D.
    cDNA Library Arrayed by: Greg Lennon, Ph.D.
    DNA Sequencing by: Washington University Genome Sequencing Center
    Clone distribution: NCI-CGAP clone distribution information can be
    found through the I.M.A.G.E. Consortium/LLNL at:
    www-bio.llnl.gov/bbrp/image/image.html
    Seq primer: -40ml3 fwd. RT from Amersham
    High quality sequence stop: 288.
  FEATURES
    Location/Qualifiers
    1. .307
      /organism="Homo sapiens"
      /mol_type="mRNA"
      /db_xref="taxon:9606"
      /clone="IMAGE:1417861"
      /tissue_type="carcinoid"
      /lab_host="DH10B"
      /clone_lib="NCI_CGAP_Lu5"
      /note="Organ: lung; Vector: pT7T3D-Pac (Pharmacia) with a
      modified polylinker; 1st strand cDNA was prepared from
      neuroendocrine lung carcinoid, and was then primed with a
      Not I - oligo(dT) primer. Double-stranded cDNA was ligated
      to Eco RI adaptors (Pharmacia), digested with Not I and
      cloned into the Not I and Eco RI sites of the modified
      pT7T3 vector. Library is normalized. Library was
      constructed by Bento Soares and M. Fatima Bonaldo. "
  BASE COUNT    116 a    62 c    42 g    87 t
  ORIGIN

  Query Match      84.8%; Score 17.8; DB 9; Length 307;
  Best Local Similarity 90.5%; Pred. No. 4.6e+02;
  Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

  QY  1 AAGCTGATTGATGACACAGG 21
      |||||
  Db   247 AAACCTGGTTGATGACACAGG 227

  RESULT 4
  AI572687/c
  LOCUS
  DEFINITION
    tr78b10.x1 NCI_CGAP_Pan1 Homo sapiens cDNA clone IMAGE:2224411 3',
    mRNA sequence.
  ACCESSION
    AI572687
  VERSION
    AI572687.1 GI:4536061
  KEYWORDS
    EST.
  SOURCE
    Homo sapiens (human)
  ORGANISM
    Homo sapiens
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
  1 (bases 1 to 339)
  NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
  National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
  TITLE

```

Tumor Gene Index

JOURNAL

COMMENT

Unpublished
Contact: Robert Strausberg, Ph.D.

Email: csapsb@mail.nih.gov

Life Technologies catalog #: 11548-013

DNA sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/bbrp/image/image.html

Insert Length: 1905 Std Error: 0.00

Seq primer: -400P from Gibco

High quality sequence stop: 332

POLYA-No.

FEATURES

source

Location/Qualifiers

1..339

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clones="IMAGE:222441"

/tissue="IMAGE:adenocarcinoma"

/lab_host="DH10B"

/clone_lib="NCI CGAP_Pan1"

/note="Organ: Pancreas; Vector: pCMV-SPORT6; Site 1: SalI;

Site 2: NotI; Cloned unidirectionally. Primer: Oligo dt.

Average insert size 1.72 kb. Life Technologies catalog #:

11548-013"

BASE COUNT

ORIGIN

129 a

67 c

46 g

97 t

Query Match

Best Local Similarity 84.8%; Score 17.8; DB 9; Length 339;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy

1 AAGCTGATTGATGACCAGG 21

|||||

Db 253 AAACCTGGTGTGACCAGG 233

RESULT 5

BH233391

LOCUS

BH233391 BH233391 380 bp DNA linear GSS 03-DEC-2001

CH230-53J16.TU CHORI-230 Segment 1 Rattus norvegicus genomic clone

CH230-53J16, genomic survey sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Rattus norvegicus (Norway rat)

Rattus norvegicus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;

Rattus.

1 (bases 1 to 380)

Zhao, S., Shetty, J., Shatsman, S., Tseng, G., Geer, K., Shvartsbeyn

A., Gebregorgis, E., Overton, L., Russell, D., Chen, D., Riggs, F., de

Jong, P., and Fraser, C.M.

Rat BAC End Sequences from Library CHORI-230 EcoRI segment

Unpublished

Other GSSs: CH230-53J16.TV

Contact: Shaying Zhao

Department of Eukaryotic Genomics

The Institute for Genomic Research

9712 Medical Center Dr., Rockville, MD 20850, USA

Tel: 301 838 0200

Fax: 301 838 0208

Email: szhao@tigr.org

Clones are derived from the rat BAC library CHORI-230

(http://www.chori.org/bacpac/rat230.html). For BAC library

availability, please contact Pieter de Jong (pdejong@mail.cho.org).

Clones may be purchased from BACPAC Resources

(http://www.chori.org/bacpac/or ering_information.html). BAC end

page: http://www.tigr.org/cdb/bac_ends/rat/bac_end_inro.html

Plate: 53 row: J column: 16

Seq primer: SP6

Class: BAC ends.

FEATURES

source

Location/Qualifiers

1..380

/organism="Rattus norvegicus"

/mol_type="genomic DNA"

/strain="BN/SeNHsd/MCW"

/db_xref="taxon:10116"

/clone="CH230-53J16"

/sex="Female"

/cell_type="Brain"

/clone_lib="CHORI-230 Segment 1"

/note="Vector: pTARAC2.1; Site 1: EcoRI; Site 2: EcoRI;

CHORI-230 Rat (BN/SeNHsd/MCW) BAC library produced by

Pieter de Jong"

BASE COUNT 105 a

89 c

103 g

83 t

Query Match

Best Local Similarity 84.8%; Score 17.8; DB 28; Length 380;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy

1 AAGCTGATTGATGACCAGG 21

|||||

Db 147 AAGCTGATTGATGATCAGAG 167

RESULT 6

CA941354/c

LOCUS

CA941354 1r34b01.x1 HR85 islet Homo sapiens cDNA clone IMAGE:554961 3',

mRNA sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Homo sapiens

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

1 (bases 1 to 545)

Melton, D., Brown, J., Kenty, G., Permutt, A., Lee, C., Kaestner, K.,

Lenishka, I., Searce, M., Brestelli, J., Gradwohl, G., Clifton, S.,

Hillier, L., Marra, M., Pape, D., Wyllie, T., Martin, J., Blistain, A.,

Schmitt, A., Theising, B., Ritter, E., Ronko, I., Bennett, J., Cardenas

M., Gibbons, M., McCann, R., Cole, R., Tsagareishvili, R., Williams, T.

, Jackson, Y. and Bowers, Y.

Endocrine Pancreas Consortium

Unpublished

Contact: Douglas Melton, Klaus H. Kaestner, & Hiroshi Inoue

Endocrine Pancreas Consortium

Harvard University, Howard Hughes Medical Institute

Dept of Molecular and Cellular Biology, 7 Divinity Ave, Cambridge,

MA 02138

Tel: 617-495-1812

Fax: 617-495-8557

Email: dmelton@biohpc.harvard.edu

Library was constructed by Dr. Hiroshi Inoue DNA sequencing by:

Washington University Genome Sequencing Center For information on

obtaining a clone please contact: Dr. Hiroshi Inoue

(hinoue@im.wustl.edu)

Seq primer: -40RP from Gibco

High quality sequence stop: 452.

Location/Qualifiers

1..545

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clones="IMAGE:554961"

/tissue="Purified pancreatic islet"

/lab_host="DH10B"

/clone_lib="HR85 islet"

/note="Organ: Pancreas; Vector: pBluescript SK(-); Site 1:

NotI; Site 2: XhoI; cDNA made by oligo-dT priming.

Size-selected on agarose gel. Average insert size ~1kb. 5'

XhoI site was destroyed after directional cloning.
 Amplified once. Contact information: Hiroshi Inoue, MD,
 Metabolism Div. (Alan Permutt Lab), Washington University
 School of Medicine, Box 8127, 660 South Euclid Ave., St.
 Louis, MO 63110, E-mail: hinoue@ingate.wustl.edu, Tel:
 314-362-1916, Fax: 314-747-2692."

BASE COUNT 184 a 100 c 88 g 173 t

ORIGIN

Query Match 84.8%; Score 17.8; DB 14; Length 545;
 Best Local Similarity 90.5%; Pred. No. 6e+02; Indels 0; Gaps 0;
 Matches 19; Conservative 0; Mismatches 2;

QY 1 AAGCTTGATTGATGACCAGG 21
 |||||
 Db 257 AAACCTGGTTGATGACCAGG 237

RESULT 7
 A1816423/c
 LOCUS
 DEFINITION au47907.x1 Schneider fetal brain 00004 Homo sapiens cDNA clone
 IMAGE:2517948 3', mRNA sequence.
 ACCESSION A1816423.1 GI:5431969
 VERSION EST.
 KEYWORDS Homo sapiens (human)
 SOURCE
 ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 592)
 Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 Hillier, L., Allen, M., Bowles, L., Dubuque, T., Geisel, G., Jost, S.,
 Krizman, D., Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin
 J., Moore, B., Schellenberg, K., Steptoe, M., Tan, P., Theising, B.,
 White, Y., Wylie, T., Waterston, R. and Wilson, R.
 WashU-NCI human EST Project
 Unpublished
 Contact: Wilson RK
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: est@watson.wustl.edu
 This clone is available royalty-free through LInL; contact the
 IMAGE Consortium (info@image.llnl.gov) for further information.
 Seq primer: -40UP from Gibco
 High quality sequence stop: 469.

FEATURES source
 1..592
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:2517948"
 /sex="male"
 /tissue_type="frontal lobe"
 /dev_stage="5 months post-conception"
 /lab_host="DH10B"
 /clone_lib="Schneider fetal brain 00004"
 /note="Organ: brain; Vector: pBluescript SK (Stratagene);
 Site 1: S81; Site 2: XhoI; Double-stranded cDNA was
 prepared from human fetal brain tissue. 5' and 3'
 adaptors were used in cloning as follows: 5' adaptor
 sequence:
 5'-GAGAGAGAGAGAGCTCAAGGATCCTTAATTAATTAATCCCGCCCCCCC-3'
 and 3' adaptor sequence:
 5'-GAGAGAGAGAGAGCTCGTTTCTTTTCTTTT-3'. The library was
 size-selected for >0.5 Kb inserts and has an average
 insert size estimated at 1.2 Kb. This library was
 constructed using the CAP-trapper method for full-length
 enrichment and has not undergone amplification. Library
 was constructed by Dr. Claudio Schneider (LNCIB-Area
 Science Park, Trieste, Italy)."

BASE COUNT 195 a 106 c 98 g 192 t 1 others

ORIGIN

Query Match 84.8%; Score 17.8; DB 9; Length 592;
 Best Local Similarity 90.5%; Pred. No. 6.2e+02; Indels 0; Gaps 0;
 Matches 19; Conservative 0; Mismatches 2;

QY 1 AAGCTTGATTGATGACCAGG 21
 |||||
 Db 259 AAACCTGGTTGATGACCAGG 239

RESULT 8
 AG121419/c
 LOCUS
 DEFINITION Pan troglodytes DNA, clone: PTB-130H10.F, genomic survey sequence.
 AG121419
 ACCESSION AG121419.1 GI:16650584
 VERSION GSS.
 KEYWORDS Pan troglodytes (chimpanzee)
 SOURCE
 ORGANISM Pan troglodytes

REFERENCE 1
 Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Pan.
 Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T.D., Yada, T.,
 Totoki, Y., Watanabe, H. and Sakaki, Y.
 BAC end sequences of Library PTB
 Unpublished
 2 (bases 1 to 671)
 Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T.D., Yada, T.,
 Totoki, Y., Watanabe, H. and Sakaki, Y.
 Direct Submission
 Submitted (02-AUG-2001) Asao Fujiyama, The Institute of Physical
 and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
 1-7-22 Suehiro-chou, Tsukumi-ku, Yokohama, Kanagawa 230-0045, Japan
 (E-mail:chimbases@sc.riken.go.jp, URL:http://bgp.gsc.riken.go.jp/,
 tel:81-45-503-9111, Fax:81-45-503-9170)
 Clones are derived from the chimpanzee BAC library PTB This BAC end
 was generated during the R&D process and may have higher chance of
 clone tracking errors.

COMMENT

PRIMERS
 Sequencing: -21M13
 LIBRARY
 Vector : pKS145
 R.Site 1 : SacI
 R.Site 2 : SacI.
 Location/Qualifiers
 1..671
 /organism="Pan troglodytes"
 /mol_type="genomic DNA"
 /db_xref="taxon:9598"
 /clone="PTB-130H10.F"
 /sex="male"
 /cell_type="lymphoblast"
 /clone_lib="PTB Chimpanzee Male BAC Library"
 BASE COUNT 195 a 161 c 152 g 163 t

ORIGIN

Query Match 84.8%; Score 17.8; DB 29; Length 671;
 Best Local Similarity 90.5%; Pred. No. 6.6e+02; Indels 0; Gaps 0;
 Matches 19; Conservative 0; Mismatches 2;

QY 1 AAGCTTGATTGATGACCAGG 21
 |||||
 Db 414 AAGCTTGATTGATGACCAGG 394

RESULT 9
 AG186976/c
 LOCUS
 DEFINITION Pan troglodytes DNA, clone: PTB-004E01.R, genomic survey sequence.
 AG186976
 ACCESSION AG186976
 VERSION AG186976.1 GI:18149509
 KEYWORDS GSS.

SOURCE Pan troglodytes (chimpanzee)
 ORGANISM Pan troglodytes
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.

REFERENCE 1
 AUTHORS Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T.D., Yada, T., Totoki, Y., Watanabe, H. and Sakaki, Y.
 TITLE BAC end sequences of Library PTB
 JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 708)
 AUTHORS Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T.D., Yada, T., Totoki, Y., Watanabe, H. and Sakaki, Y.
 TITLE Direct Submission
 JOURNAL Submitted (08-DEC-2001) Asao Fujiyama, The Institute of Physical and Chemical Research (RIKEN), Genomic Sciences Center (GSC); 1-7-22 Suehiro-chou, Tsukumi-ku, Yokohama, Kanagawa 230-0045, Japan (E-mail: chibbes@gs.c.riken.go.jp, URL: http://hgp.gsc.riken.go.jp/, Tel: 81-45-503-9111, Fax: 81-45-503-9170)

COMMENT Clones are derived from the chimpanzee BAC library PTB. This BAC end was generated during the R&D process and may have higher chance of clone tracking errors.

PRIMERS
 Sequencing: M13Rev
 LIBRARY
 Vector : PKS145
 R.Site 1 : SacI
 R.Site 2 : SacI
 Location/Qualifiers
 1..708
 /organism="Pan troglodytes"
 /mol_type="genomic DNA"
 /db_xref="taxon:9598"
 /clone="PTB-094E01.R"
 /sex="male"
 /cell_type="lymphoblast"
 /clone_lib="PTB Chimpanzee Male BAC Library"
 201 a 165 c 156 g 182 t 3 others

BASE COUNT
 ORIGIN

Query Match 84.8%; Score 17.8; DB 29; Length 708;
 Best Local Similarity 90.5%; Pred. No. 6.7e+02;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 AAGCTGATTCATGACCCAGG 21
 Db 398 AAGCTGATTCATGACCCAGG 378

RESULT 10
 LOCUS BU245702 913 bp mRNA linear EST 26-NOV-2002
 DEFINITION 603783316F1 CSEQCHN34 Gallus gallus cdna clone CHEST73504 5', mRNA sequence.
 ACCESSION BU245702
 VERSION BU245702.1 GI:25493069
 EST
 KEYWORDS Gallus gallus (chicken)
 SOURCE Gallus gallus
 ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Archosauria; Aves; Neognathae; Galliformes; Phasianidae; Phasianinae; Gallus.
 1 (bases 1 to 913)
 REFERENCE Boardman, P.E., Sanz-Ezquerro, J., Overton, I.M., Burt, D.W., Bosch, E., Fong, W.T., Tickle, C., Brown, W.R.A., Wilson, S.A. and Hubbard, S.J.
 AUTHORS A Comprehensive Collection of Chicken cDNAs
 TITLE Curr. Biol. 12 (22), 1965-1969 (2002)
 JOURNAL 22335534
 MEDLINE 22335534
 PUBMED 12445392
 COMMENT Contact: Simon Hubbard
 Department of Biomolecular Sciences
 University of Manchester Institute of Science and Technology (UMIST)
 PO Box 88, Manchester, M60 1QD, UK

Tel: 01612008930
 Fax: 01612360409
 Email: Simon.Hubbard@umist.ac.uk.

FEATURES
 source
 Location/Qualifiers
 1..913
 /organism="Gallus gallus"
 /mol_type="mRNA"
 /strain="Compton Line 151"
 /db_xref="taxon:9031"
 /clone="CHEST73504"
 /sex="Female"
 /dev_stage="adult"
 /lab_host="DH10B"
 /clone_lib="CSEQCHN34"
 /note="Organ: liver; Vector: pBluescript II KS(+); Site_1: EcoRI; Site_2: NotI; This normalized library was constructed from 1 million independent clones. cDNA synthesis was initiated using an oligo(dT) primer, using methylated C in the first strand synthesis reaction. Following this first strand reaction, double-stranded cDNA was blunted, ligated to NotI adapters, digested with EcoRI, size-selected, and cloned into the NotI and EcoRI compatible sites of a custom modified MCS of the pBluescript (KS+) vector. The library was normalized in 2 rounds using conditions adapted from Soares et al., PNAS (1994) 91: 9228-9232 and Bonaldo et al., Genome Research 6 (1996): 791, except that a significantly longer reannealing hybridization was used."

BASE COUNT 217 a 230 c 207 g 259 t

ORIGIN

Query Match 84.8%; Score 17.8; DB 13; Length 913;
 Best Local Similarity 90.5%; Pred. No. 7.6e+02;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 AAGCTGATTCATGACCCAGG 21
 Db 145 AAGCTGATTCATGACCCAGG 165

RESULT 11
 LOCUS BG119393 963 bp mRNA linear EST 30-JAN-2001
 DEFINITION 602349076F1 NIH_MGC_90 Homo sapiens CDNA clone IMAGE:4444074 5', mRNA sequence.
 ACCESSION BG119393
 VERSION BG119393.1 GI:12612899
 EST.
 KEYWORDS Homo sapiens (human)
 SOURCE Homo sapiens
 ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 963)
 REFERENCE NIH-MGC http://mgc.nci.nih.gov/.
 AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)
 TITLE Unpublished
 JOURNAL Contact: Robert Strausberg, Ph.D.
 COMMENT Email: cgabbs-remail.nih.gov
 Tissue Procurement: ATCC
 CDNA Library Preparation: Life Technologies, Inc.
 CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov
 Plate: LLAM10219 row: g column: 19
 High quality sequence stop: 658.

FEATURES
 source
 Location/Qualifiers
 1..963
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:4444074"

```

/tissue_type="adenocarcinoma, cell line"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NIH MGC 90"
/notes="Organ: liver; Vector: pCMV-SPORT6; Site 1: NotI;
Site 2: SalI; Cloned unidirectionally; oligo-dT primed.
Average insert size 1.7 Kb. Library enriched for
full-length clones and constructed by Life Technologies.
Note: this is a NIH MGC Library."
BASE COUNT      285 a 199 c 199 g 280 t
ORIGIN

Query Match      84.8%; Score 17.8; DB 10; Length 963;
Best Local Similarity 90.5%; Pred. No. 7.8e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAGCTGATTGATGACCCAGG 21
    |||||
Db 570 AAACCTGGTTGATGACCCAGG 590

RESULT 12
AL523235/c
LOCUS      1201 bp      mRNA      linear      EST 22-MAY-2003
DEFINITION Homo sapiens NEUROBLASTOMA COT 25-NORMALIZED Homo sapiens
cDNA clone CS0DC001YE18 3-PRIME, mRNA sequence.
ACCESSION  AL523235
VERSION     AL523235.2 GI:31041496
KEYWORDS    EST.
SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
REFERENCE   1 (bases 1 to 1201)
AUTHORS    Li, W.B., Gruber, C., Jessee, J. and Polayes, D.
TITLE      Full-length cDNA libraries and normalization
JOURNAL    Unpublished
COMMENT    On Feb 13, 2001 this sequence version replaced gi:12786728.
Contact: Genoscope
Genoscope - Centre National de Sequencage
BP 191 91006 EVRY cedex - France
Email: segref@genoscope.cns.fr, Web : www.genoscope.cns.fr
Library was constructed by Life Technologies, a division of
Invitrogen. This sequence belongs to sequence cluster 10668.r, and
it belongs to a clone representative of this cluster. For more
information about this cluster and the virtual cDNA, see
http://www.genoscope.cns.fr/
cgi-bin/cluster.cgi?seq=CS0DC001BC09NP1&cluster=10668.r. Contact :
Feng Liang Email : fliang@lifetech.com URL : Corporation 1600
http://fulllength.invitrogen.com/ Invitrogen Corporation 1600
Faraday Avenue Genoscope sequence ID : CS0DC001BC09NP1.
Location/Qualifiers
source      1. 1201
            /organism="Homo sapiens"
            /mol_type="mRNA"
            /db_xref="taxon:9606"
            /clone="CS0DC001YE18"
            /tissue_type="NEUROBLASTOMA COT 25-NORMALIZED"
            /clone_lib="Homo sapiens NEUROBLASTOMA COT 25-NORMALIZED"
            /note="First strand cDNA was primed with a NotI-oligo(dT)
primer. Five prime end enriched, double-strand cDNA was
digested with NotI and cloned into the NotI and EcoR V
sites of the pCMVSPORT 6 vector. Library was normalized."
BASE COUNT  358 a 206 c 252 g 311 t 74 others
ORIGIN

Query Match      84.8%; Score 17.8; DB 9; Length 1201;
Best Local Similarity 90.5%; Pred. No. 8.6e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAGCTGATTGATGACCCAGG 21
    |||||
Db 236 AAACCTGGTTGATGACCCAGG 216

/tissue_type="adenocarcinoma, cell line"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NIH MGC 90"
/notes="Organ: liver; Vector: pCMV-SPORT6; Site 1: NotI;
Site 2: SalI; Cloned unidirectionally; oligo-dT primed.
Average insert size 1.7 Kb. Library enriched for
full-length clones and constructed by Life Technologies.
Note: this is a NIH MGC Library."
BASE COUNT      285 a 199 c 199 g 280 t
ORIGIN

Query Match      84.8%; Score 17.8; DB 10; Length 963;
Best Local Similarity 90.5%; Pred. No. 7.8e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAGCTGATTGATGACCCAGG 21
    |||||
Db 570 AAACCTGGTTGATGACCCAGG 590

RESULT 13
AZ072191
LOCUS      565 bp      DNA      linear      GSS 31-MAR-2000
DEFINITION RPCI-23-39618.TV RPCI-23 Mus musculus genomic clone RPCI-23-39618,
genomic survey sequence.
ACCESSION  AZ072191
VERSION    AZ072191
KEYWORDS   GSS.
SOURCE     Mus musculus (house mouse)
ORGANISM   Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE   1 (bases 1 to 565)
AUTHORS     Zhao, S., Nierman, W., Feldblyum, T., Malek, J., Shatsman, S., Akinret
B., Levins, M., McGann, S., Tsegaye, G., Geer, K., Krol, M., de Jong, P.
and Fraser, C.M.
TITLE      Mouse BAC End Sequences from Library RPCI-23
JOURNAL    Unpublished
COMMENT    Contact: Shaying Zhao
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: szhao@tigr.org
Clones are derived from the mouse BAC library RPCI-23. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/orderingframe.htm)
or from Resea ch Genetics (info@resgen.com). BAC end page:
http://www.tigr.org/tdb/bac/ends/mouse/bac_end_intro.html
Plate: 396 row: I column: 8
Seq primer: T7
Class: BAC ends.
Location/Qualifiers
source      1. 565
            /organism="Mus musculus"
            /mol_type="genomic DNA"
            /strain="C57BL/6J"
            /db_xref="taxon:10090"
            /clone="RPCI-23-39618"
            /sex="Female"
            /lab_host="DH10B"
            /clone_lib="RPCI-23"
            /notes="Organ: Kidney/Brain; Vector: pBACe3.6; Site 1:
EcoRI; Site 2: EcoRI; Female C57BL/6J mouse kidney and/or
brain genomic DNA was isolated and partially digested
with a combination of EcoRI and EcoRI Methylase. Size
selected DNA was cloned into the pBACe3.6 vector at the
EcoRI sites. The ligation products were transformed into
DH10B electrocompetent cells (BRL Life Technologies)."
BASE COUNT  142 a 131 c 162 g 130 t
ORIGIN

Query Match      82.9%; Score 17.4; DB 28; Length 565;
Best Local Similarity 94.7%; Pred. No. 9.4e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 AGCCTGATTGATGACCCAGG 20
    |||||
Db 26 AGCCTGATTGATGACCCAGG 44

RESULT 14
BE729033/c
LOCUS      914 bp      mRNA      linear      EST 15-SEP-2000
DEFINITION 601561212F1 NIH MGC 20 Homo sapiens cDNA clone IMAGE:3830835 5',
mRNA sequence.
ACCESSION  BE729033
VERSION    BE729033.1 GI:10143025
KEYWORDS   EST.
SOURCE     Homo sapiens (human)

```

ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 914)
 AUTHORS NIH-MGC <http://mgi.nci.nih.gov/>
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-r@mail.nih.gov
 Tissue Procurement: ATCC/DCTD/DTT
 cDNA Library Preparation: Ling Hong/Rubin Laboratory
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at: image.llnl.gov
 Plate: LLCW505 row: h column: 04
 High quality sequence stop: 694.

FEATURES
 source
 1..914
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:3830835"
 /tissue_type="melanotic melanoma"
 /lab_host="DH10B (phage-resistant)"
 /clone_lib="NIH MGC 20"
 /note="Organ: skin; Vector: pOTB7; Site 1: XhoI; Site 2:
 EcoRI; cDNA made by oligo-dT priming. Directionally
 cloned into EcoRI/XhoI sites using the following 5'
 adaptor: GGCACGAG(G). Size-selected >500bp for average
 insert size 1.8kb. Library constructed by Ling Hong in
 the laboratory of Gerald M. Rubin (University of
 California, Berkeley) using ZAP-cDNA synthesis kit
 (Stratagene) and Superscript II RT (Life Technologies)."
 BASE COUNT 212 a 241 c 238 g 223 t
 ORIGIN

Query Match 82.9%; Score 17.4; DB 10; Length 914;
 Best Local Similarity 94.7%; Pred. No. 1.2e+03;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 AAGCTGATTGATGACCAG 19
 |||||
 Db 626 AAGCTGATTGATGACCAG 608

RESULT 15
 AI818091/c
 LOCUS AI818091 543 bp mRNA linear EST 21-DEC-1999
 DEFINITION WK27a09.x1 NCI CGAP Brn25 Homo sapiens cDNA clone IMAGE:241352B 3,
 similar to SW:HS1_FACSU 034520 ATP PHOSPHORIBOSYLTRANSFERASE 1,
 mRNA sequence.
 ACCESSION AI818091
 VERSION AI818091.1 GI:5437170
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 543)
 AUTHORS NCI/NINDS-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
 TITLE National Cancer Institute / National Institute of Neurological
 Disorders and Stroke, Brain Tumor Genome Anatomy Project
 (CGAP/ETGAP), Tumor Gene Index
 JOURNAL Unpublished
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-r@mail.nih.gov
 Tissue Procurement: David N. Louis, M.D., Myrna R. Rosenfeld M.D.,
 Ph.D.
 cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
 Bonaldo, Ph.D.
 cDNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
 Insert Length: 904 Std Error: 0.00
 Seq primer: -40UP from Gibco
 High quality sequence stop: 485.

FEATURES
 source
 Location/Qualifiers
 1..543
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:2413528"
 /tissue_type="anaplastic oligodendroglioma"
 /lab_host="DH10B"
 /clone_lib="NCI CGAP Brn25"
 /note="Organ: brain; Vector: pT7T3D-Pac (Pharmacia) with a
 modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st
 strand cDNA was primed with a Not I - oligo(dT) primer [5'
 TGTTACCAATCTGAAGTGGAGCGCGCATAGGTTTTTTTTTTTTTTTTTT
 T 3']; double-stranded cDNA was ligated to Eco RI
 adaptors (Pharmacia), digested with Not I and cloned into
 the Not I and Eco RI sites of the modified pT7T3 vector.
 Library is normalized, and was constructed by Bento
 Soares and M. Fatima Bonaldo."
 BASE COUNT 99 a 174 c 180 g 90 t
 ORIGIN

Query Match 81.0%; Score 17; DB 9; Length 543;
 Best Local Similarity 100.0%; Pred. No. 1.4e+03;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GCCTGATTGATGACCAG 19
 |||||
 Db 494 GCCTGATTGATGACCAG 478

Search completed: February 19, 2004, 23:14:36
 Job time : 1575 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: February 19, 2004, 19:59:22 ; Search time 175.5 Seconds
(without alignments)
323.010 Million cell updates/sec

Title: US-10-085-108-21_COPY_711_731

Perfect score: 21

Sequence: 1 AAGCTGATTGATGACCAGG 21

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 2552756 seqs, 1349719017 residues

Total number of hits satisfying chosen parameters: 5105512

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : N_Geneseq_19Jun03.*

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22: /SIDSI/gcgdata/geneseq/geneseq-nbml/NA2001A.DAT.*
23: /SIDSI/gcgdata/geneseq/geneseq-nbml/NA2001B.DAT.*
24: /SIDSI/gcgdata/geneseq/geneseq-nbml/NA2002.DAT.*
25: /SIDSI/gcgdata/geneseq/geneseq-nbml/NA2003.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	21	100.0	21	25	Human MAGE-C3 expr
C 2	21	100.0	425	22	Human breast cell
C 3	21	100.0	425	22	Human foetal liver
C 4	21	100.0	425	22	Probe #4696 for ge
C 5	21	100.0	425	22	Human brain expres
C 6	21	100.0	425	22	Human bone marrow
C 7	21	100.0	425	22	Probe #4815 for ge
C 8	21	100.0	425	22	Probe #4923 used t

C 9	21	100.0	425	22	AAI04666	Probe #4657 used t
C 10	21	100.0	425	23	ABS29927	Human liver single
C 11	21	100.0	425	24	ABS04866	Human genome-deriv
C 12	21	100.0	1041	25	ABS95006	cDNA encoding huma
C 13	21	100.0	7806	16	ABS88354	DNA encoding novel
C 14	17	81.0	336	23	AAI24776	Human gene signatu
C 15	16.8	80.0	504	24	ABS69805	Novel murine polyn
C 16	16.2	77.1	202	19	AAI10827	Human biallelic po
C 17	16.2	77.1	419	23	ABV17420	Human prostate exp
C 18	16.2	77.1	456	22	ABA58928	Human foetal liver
C 19	16.2	77.1	456	22	AAK07090	Human brain expres
C 20	16.2	77.1	456	22	AAK32830	Human bone marrow
C 21	16.2	77.1	456	22	AAI38645	Probe #7331 used t
C 22	16.2	77.1	456	23	ABS32555	Human liver single
C 23	16.2	77.1	456	24	ABS07634	Human genome-deriv
C 24	16.2	77.1	458	23	ABV47214	Human prostate exp
C 25	16.2	77.1	964	22	ABA15191	Human nervous syst
C 26	16.2	77.1	1236	23	ABL03601	Drosophila melanog
C 27	16.2	77.1	1781	22	ABAI18978	Human nervous syst
C 28	16.2	77.1	1920	22	ABAI10827	Manihot esculenta
C 29	16.2	77.1	2204	22	ABA15194	Human nervous syst
C 30	16.2	77.1	3171	22	ACQ85455	Cold-active beta g
C 31	16.2	77.1	3479	23	ABL03600	Drosophila melanog
C 32	16.2	77.1	3968	23	ABL03602	Drosophila melanog
C 33	16.2	77.1	4912	21	AAA65349	NDO related comple
C 34	16.2	77.1	6779	21	AAA65350	NDO related comple
C 35	16.2	77.1	7325	24	AA894817	Human DNA sequence
C 36	15.8	75.2	470	22	ABA26707	Probe #5173 for ge
C 37	15.8	75.2	1378	23	ABL07577	Drosophila melanog
C 38	15.8	75.2	3116	23	ABL03310	Drosophila melanog
C 39	15.8	75.2	3378	23	ABL07576	Drosophila melanog
C 40	15.8	75.2	4764	23	ABL05675	Drosophila melanog
C 41	15.8	75.2	6147	22	ABAI14804	Human nervous syst
C 42	15.8	75.2	7300	23	ABL05674	Drosophila melanog
C 43	15.4	73.3	312	22	AAH66909	C glutamic codin
C 44	15.4	73.3	320	22	AAK87704	Human digestive sy
C 45	15.4	73.3	320	22	AAS31687	Human liver associ

ALIGNMENTS

RESULT 1

ABX95022/c

ID ABX95022 standard; DNA; 21 BP.

XX ABX95022;

AC ABX95022;

XX

DT 05-JUN-2003 (first entry)

XX Human MAGE-C3 expression pattern anlaysis RT-PCR antisense primer.

DE TRAP; ss; tumour rejection antigen precursor; cytolytic T-cell; CTL;

XX tumour; seminoma; bladder transitional-cell carcinoma; NSCLC; adaptor;

KW head-and-neck squamous-cell carcinoma; breast carcinoma; sarcoma;

KW cutaneous melanoma; nonsmall cell lung cancer; RT-PCR; primer; MAGE-C3;

KW human; reverse transcription.

XX Homo sapiens.

OS US2002176865-A1.

FN 28-NOV-2002.

PD 01-MAR-2002; 2002US-0085108.

XX 09-FEB-2000; 2000US-0501104.

PR 25-APR-1997; 97US-0845528.

PR 24-APR-1998; 98US-0066281.

PR 17-DEC-1999; 99US-0468433.

XX (LUCAS/)

PA (BOON/)

PA BOON-FALLEUR T.

XX Lucas S, Boon-Falleur T;
 XX WPI; 2003-328468/31.
 XX
 XX Novel isolated nucleic acid encoding tumor rejection antigen precursor
 PT MAGE-C3, MAGE-B5, or MAGE-B6, useful as diagnostic probes to determine
 PT presence of abnormal e.g., tumor cells expressing MAGE-C1, MAGE-B5 or
 PT MAGE-B6
 XX
 XX Example 12; Page 13; 59pp; English.
 XX
 XX The invention relates to an isolated nucleic acid molecule which encodes
 CC a tumour rejection antigen precursor (TRAP) having an amino acid sequence
 CC of a TRAP encoded by a fully defined MAGE-C3, MAGE-B5, or MAGE-B6
 CC polynucleotide sequence. Also disclosed is a method which is useful for
 CC determining presence of cytolytic T-cells specific for complexes of human
 CC leukocyte antigen (HLA) and a peptide derived from the nucleic acid in a
 CC cytotoxic T-lymphocyte (CTL)-containing sample. The nucleic acid is
 CC useful as a diagnostic probe to determine the presence of abnormal
 CC (tumour) cells such as seminoma, bladder transitional-cell carcinoma,
 CC head-and-neck squamous-cell carcinoma, breast carcinoma, sarcoma,
 CC cutaneous melanoma or non-small cell lung cancer (NSCLC) which express
 CC MAGE-C1, MAGE-B5 or MAGE-B6. The nucleic acid is useful for diagnosing a
 CC disorder characterised by expression of MAGE-C1, MAGE-B5 or MAGE-B6 TRAPS
 CC or tumour rejection antigens (TRAs). The present sequence represents the
 CC human MAGE-C3 expression pattern analysis reverse transcription (RT)-PCR
 CC antisense primer.
 XX
 XX Sequence 21 BP; 4 A; 7 C; 4 G; 6 T; 0 other;
 SQ
 Query Match 100.0%; Score 21; DB 25; Length 21;
 Best Local Similarity 100.0%; Pred. No. 0.39;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 AGCCTGATTGATGACACGGG 21
 Db |||||
 21 AGCCTGATTGATGACACGGG 1
 RESULT 2
 ABA46075/c
 ID ABA46075 standard; DNA; 425 BP.
 XX
 AC ABA46075;
 XX
 DT 01-FEB-2002 (first entry)
 XX
 DE Human breast cell single exon nucleic acid probe #4770.
 XX
 KW Human; microarray; single exon probe; gene expression; breast;
 KW disease; cancer; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200157271-A2.
 XX
 PD 09-AUG-2001.
 XX
 PF 30-JAN-2001; 2001WO-US00662.
 XX
 PR 04-FEB-2000; 2000US-0180312.
 XX
 PR 26-MAY-2000; 2000US-0207456.
 XX
 PR 30-JUN-2000; 2000US-0608408.
 XX
 PR 03-AUG-2000; 2000US-0632366.
 XX
 PR 21-SEP-2000; 2000US-0234687.
 XX
 PR 27-SEP-2000; 2000US-0236359.
 XX
 PR 04-OCT-2000; 2000GB-0024263.
 XX
 PA (MOLE-) MOLECULAR DYNAMICS INC.
 XX
 PI Penn SG, Hanzel DK, Chen W, Rank DR;
 XX
 PI WPI; 2001-483447/52.
 XX
 XX Human genome-derived single exon nucleic acid probes useful for

DR WPI; 2001-496933/54.
 XX
 XX New spatially-addressable set of single exon nucleic acid probes,
 PT useful for measuring gene expression in sample derived from human
 PT breast, comprises number of single exon nucleic acid probes
 XX
 XX Claim 1; SEQ ID NO 4770; 327pp + sequence listing; English.
 XX
 XX The invention relates to a spatially-addressable set of single exon
 CC nucleic acid probes for measuring gene expression in a sample derived
 CC from human breast and BT 474 cells. The method involves contacting
 CC the probes with a collection of detectably labelled nucleic acids
 CC derived from mRNA of human breast, and then measuring the label
 CC bound to each probe of the microarray. The probes are useful for
 CC verifying the expression of regions of genomic DNA predicted to
 CC encode proteins. They are useful for gene discovery, and for
 CC determining predisposition and/or prognosing breast disease. Gene
 CC expression analysis is useful for assessing the toxicity of chemical
 CC agents on cells. The microarray of this invention presents a far greater
 CC diversity of probes for measuring gene expression, with far less bias
 CC than expressed sequence tag microarrays. The method is suitable for
 CC rapid production of functional information from genomic sequence. The
 CC present sequence is a single exon nucleic acid probe of the invention.
 CC Note: The sequence data for this patent did not form part of the
 CC printed specification, but was obtained in electronic format directly
 CC from WIPO at ftp.wipo.int/pub/published_pct_sequences.
 XX
 XX Sequence 425 BP; 112 A; 107 C; 94 G; 112 T; 0 other;
 SQ
 Query Match 100.0%; Score 21; DB 22; Length 425;
 Best Local Similarity 100.0%; Pred. No. 0.69;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 AAGCCTGATTGATGACACGGG 21
 Db |||||
 150 AAGCCTGATTGATGACACGGG 130
 RESULT 3
 ABA56617/c
 ID ABA56617 standard; DNA; 425 BP.
 XX
 AC ABA56617;
 XX
 DT 01-FEB-2002 (first entry)
 XX
 DE Human foetal liver single exon nucleic acid probe #4922.
 XX
 KW Human; foetal liver; gene expression; single exon nucleic acid probe; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200157277-A2.
 XX
 PD 09-AUG-2001.
 XX
 PF 30-JAN-2001; 2001WO-US00669.
 XX
 PR 04-FEB-2000; 2000US-0180312.
 XX
 PR 26-MAY-2000; 2000US-0207456.
 XX
 PR 30-JUN-2000; 2000US-0608408.
 XX
 PR 03-AUG-2000; 2000US-0632366.
 XX
 PR 21-SEP-2000; 2000US-0234687.
 XX
 PR 27-SEP-2000; 2000US-0236359.
 XX
 PR 04-OCT-2000; 2000GB-0024263.
 XX
 PA (MOLE-) MOLECULAR DYNAMICS INC.
 XX
 PI Penn SG, Hanzel DK, Chen W, Rank DR;
 XX
 PI WPI; 2001-483447/52.
 XX
 XX Human genome-derived single exon nucleic acid probes useful for

analyzing gene expression in human fetal liver -
Claim 1; SEQ ID NO 4922; 639pp + sequence listing; English.
The invention relates to a single exon nucleic acid probe for measuring human gene expression in a sample derived from human foetal liver. The single exon nucleic acid probes may be used for predicting, measuring and displaying gene expression in samples derived from human foetal liver. The present sequence is a single exon nucleic acid probe of the invention.
Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/published_pct_sequences.
Sequence 425 BP; 112 A; 107 C; 94 G; 112 T; 0 other;
Query Match 100.0%; Score 21; DB 22; Length 425;
Best Local Similarity 100.0%; Pred. No. 0.69; Indels 0; Gaps 0;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AAGCCTGATTGATGACCAGGG 21
DB 150 AAGCCTGATTGATGACCAGGG 130
RESULT 4
ID ABA26230/c
XZ ABA26230 standard; DNA; 425 BP.
AC ABA26230;
XX
DT 23-JAN-2002 (first entry)
DE Probe #4696 for gene expression analysis in human heart cell sample.
KW Human; gene expression; heart; microarray; vascular system; probe;
KW cardiovascular disease; hypertension; cardiac arrhythmia;
KW congenital heart disease; ss.
OS Homo sapiens.
XX
PN WO200157274-A2.
XX
PD 09-AUG-2001.
XX
PF 30-JAN-2001; 2001WO-US00666.
XX
PR 04-FEB-2000; 2000US-0180312.
PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
WPI; 2001-488899/53.
XX
PT Single exon nucleic acid probes for analyzing gene expression in human hearts -
PS Claim 1; SEQ ID NO 4696; 530pp; English.
XX
CC The present invention relates to single exon nucleic acid probes for measuring human gene expression in a sample derived from human heart. The present sequence is one such probe. The probes may be used for predicting, measuring and displaying gene expression in samples derived from the human heart via microarrays. By measuring gene expression, the probes are useful for predicting, diagnosing, grading, staging, monitoring and prognosing diseases of the human heart and vascular system

e.g. cardiovascular disease, hypertension, cardiac arrhythmias and congenital heart disease.
Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/published_pct_sequences.
Sequence 425 BP; 112 A; 107 C; 94 G; 112 T; 0 other;
Query Match 100.0%; Score 21; DB 22; Length 425;
Best Local Similarity 100.0%; Pred. No. 0.69; Indels 0; Gaps 0;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AAGCCTGATTGATGACCAGGG 21
DB 150 AAGCCTGATTGATGACCAGGG 130
RESULT 5
ID AAK04758/c
XZ AAK04758 standard; DNA; 425 BP.
AC AAK04758;
XX
DT 05-NOV-2001 (first entry)
DE Human brain expressed single exon probe SEQ ID NO: 4749.
KW Human; brain expressed exon; gene expression analysis; probe;
KW microarray; Alzheimer's disease; multiple sclerosis; schizophrenia; epilepsy; cancer; ss.
OS Homo sapiens.
XX
PN WO200157275-A2.
XX
PD 09-AUG-2001.
XX
PF 30-JAN-2001; 2001WO-US00667.
XX
PR 04-FEB-2000; 2000US-0180312.
PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
WPI; 2001-483446/52.
XX
PT Single exon nucleic acid probes for analyzing gene expression in human brains -
PS Example 4; SEQ ID NO: 4749; 650pp + Sequence Listing; English.
XX
CC The present invention provides a number of single exon nucleic acid probes which are derived from genomic sequences expressed in the human brain. They can be used to measure gene expression in brain cell samples, which may enable the diagnosis and improved treatment of nervous system diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia, epilepsy and cancers. The present sequence is one of the probes of the invention.
XX
SQ Sequence 425 BP; 112 A; 107 C; 94 G; 112 T; 0 other;
Query Match 100.0%; Score 21; DB 22; Length 425;
Best Local Similarity 100.0%; Pred. No. 0.69; Indels 0; Gaps 0;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AAGCCTGATTGATGACCAGGG 21

Db 150 AAGCCTGATTGATGACCAGG 130
|||||

RESULT 6

AAK30282/c
ID AAK30282 standard; DNA; 425 BP.

XX AC AAK30282;

XX DT 06-NOV-2001 (first entry)

XX DE Human bone marrow expressed single exon probe SEQ ID NO: 4839.
XX KW Human; bone marrow expressed exon; gene expression analysis; probe;
XX KW microarray; cancer; leukaemia; lymphoma; myeloma; ss.
XX OS Homo sapiens.

XX PN WO200157276-A2.

XX PD 09-AUG-2001.

XX PF 30-JAN-2001; 2001WO-US006568.

XX PR 04-FEB-2000; 2000US-0180312.

XX PR 26-MAY-2000; 2000US-0207456.

XX PR 30-JUN-2000; 2000US-0608408.

XX PR 03-AUG-2000; 2000US-0632366.

XX PR 21-SEP-2000; 2000US-0234687.

XX PR 27-SEP-2000; 2000US-0236359.

XX PR 04-OCT-2000; 2000GB-0024263.

XX PA (MOLE-) MOLECULAR DYNAMICS INC.

XX PI Penn SG, Hanzel DK, Chen W, Rank DR;

XX PD WPI; 2001-488901/53.

XX PT Human genome-derived single exon nucleic acid probes useful for
analyzing gene expression in human bone marrow -

XX PS Example 4; SEQ ID NO: 4839; 658pp + Sequence Listing; English.

XX CC The present invention provides a number of single exon nucleic acid
probes which are derived from genomic sequences expressed in the human
bone marrow. They can be used to measure gene expression in bone marrow
samples, which may enable the improved diagnosis and treatment of cancers
such as lymphoma, leukaemia and myeloma. The present sequence is one of
the probes of the invention.

XX SQ Sequence 425 BP; 112 A; 107 C; 94 G; 112 T; 0 other;

Query Match 100.0%; Score 21; DB 22; Length 425;

Best Local Similarity 100.0%; Pred. NO. 0.69;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AAGCCTGATTGATGACCAGG 21
|||||

Db 150 AAGCCTGATTGATGACCAGG 130
|||||

RESULT 7

AAI14882/c

ID AAI14882 standard; DNA; 425 BP.

XX AC AAI14882;

XX DT 12-OCT-2001 (first entry)

XX DE Probe #4815 for gene expression analysis in human cervical cell sample.
XX KW Probe; human; microarray; gene expression; cervical epithelial cell;

KW cervical cancer; ss.

OS Homo sapiens.

XX PN WO200157278-A2.

XX PD 09-AUG-2001.

XX PF 30-JAN-2001; 2001WO-US006670.

XX PR 04-FEB-2000; 2000US-0180312.

XX PR 26-MAY-2000; 2000US-0207456.

XX PR 30-JUN-2000; 2000US-0608408.

XX PR 03-AUG-2000; 2000US-0632366.

XX PR 21-SEP-2000; 2000US-0234687.

XX PR 27-SEP-2000; 2000US-0236359.

XX PR 04-OCT-2000; 2000GB-0024263.

XX PA (MOLE-) MOLECULAR DYNAMICS INC.

XX PI Penn SG, Hanzel DK, Chen W, Rank DR;

XX PD WPI; 2001-488901/53.

XX PT Human genome-derived single exon nucleic acid probes useful for
analyzing gene expression in human cervical epithelial cells -

XX PS Claim 25; SEQ ID NO 4815; 487pp; English.

XX CC The present invention relates to human single exon nucleic acid probes
(SENP). The present sequence is one such probe. The SENPs are derived
from human Hela cells. The SENPs can be used to produce a single exon
microarray, which can be used for measuring human gene expression in a
sample derived from human cervical epithelial cells. By measuring gene
expression, the probes are therefore useful in grading and/or staging
of diseases of the cervix, notably cervical cancer.

XX CC Note: The sequence data for this patent did not form part of the printed
specification, but was obtained in electronic format directly from WIPO
at ftp.wipo.int/pub/published_pct_sequences.

XX SQ Sequence 425 BP; 112 A; 107 C; 94 G; 112 T; 0 other;

Query Match 100.0%; Score 21; DB 22; Length 425;

Best Local Similarity 100.0%; Pred. NO. 0.69;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AAGCCTGATTGATGACCAGG 21
|||||

Db 150 AAGCCTGATTGATGACCAGG 130
|||||

RESULT 8

AAI36237/c

ID AAI36237 standard; DNA; 425 BP.

XX AC AAI36237;

XX DT 17-OCT-2001 (first entry)

XX DE Probe #4923 used to measure gene expression in human placenta sample.
XX KW Probe; microarray; human; placenta; antenatal diagnosis;
XX KW genetic disorder; ss.

XX OS Homo sapiens.

XX PN WO200157272-A2.

XX PD 09-AUG-2001.

XX PF 30-JAN-2001; 2001WO-US006663.

XX PR 04-FEB-2000; 2000US-0180312.

PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX (MOLE-) MOLECULAR DYNAMICS INC.
XX Penn SG, Hanzel DK, Chen W, Rank DR;
XX WPI; 2001-488897/53.
XX Human genome-derived single exon nucleic acid probes useful for
PT analysing gene expression in human placenta -
XX
PS Claim 25; SEQ ID No 4923; 654pp; English.
XX
XX The present invention relates to single exon nucleic acid probes (SENP).
CC The present sequence is one such probe. The probes are useful for
CC producing a microarray for predicting, measuring and displaying gene
CC expression in samples derived from human placenta. The probes are useful
CC for antenatal diagnosis of human genetic disorders.
XX
SQ Sequence 425 BP; 112 A; 107 C; 94 G; 112 T; 0 other;
Query Match 100.0%; Score 21; DB 22; Length 425;
Best Local Similarity 100.0%; Pred. No. 0.69;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AAGCTGATTGATGACACGAGG 21
DB 150 AAGCTGATTGATGACACGAGG 130
RESULT 9
AAI04666/c
ID AAI04666 standard; DNA; 425 BP.
XX
AC AAI04666;
XX
DT 09-OCT-2001 (first entry)
XX
DE Probe #4657 used to measure gene expression in human breast sample.
XX
KW Probe; human; breast disease; breast cancer; development disorder; ss;
KW inflammatory disease; proliferative breast disease; non-carcinoma tumour.
XX
OS Homo sapiens.
XX
PN WO200157270-A2.
XX
PD 09-AUG-2001.
XX
PF 29-JAN-2001; 2001WO-US00661.
XX
PR 04-FEB-2000; 2000US-0180312.
PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX Penn SG, Hanzel DK, Chen W, Rank DR;
XX WPI; 2001-476286/51.
XX Novel single exon nucleic acid probe used to measuring gene expression
PT in a human breast -
XX

PS Claim 25; SEQ ID No 4657; 322pp; English.
XX
XX The present invention relates to novel single exon nucleic acid probes.
CC The present sequence is one such probe. The probes are useful for
CC measuring human gene expression in a human breast sample, where the probe
CC hybridises at high stringency to a nucleic acid expressed in the human
CC breast. The probes are useful for predicting, diagnosing, grading,
CC staging, monitoring and prognosing diseases of the human breast,
CC particularly those diseases with polygenic aetiology. The diseases
CC include: breast cancer, disorders of development, inflammatory diseases
CC of the breast, fibrocystic changes, proliferative breast disease and
CC non-carcinoma tumours.
CC Note: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 425 BP; 112 A; 107 C; 94 G; 112 T; 0 other;
Query Match 100.0%; Score 21; DB 22; Length 425;
Best Local Similarity 100.0%; Pred. No. 0.69;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AAGCTGATTGATGACACGAGG 21
DB 150 AAGCTGATTGATGACACGAGG 130
RESULT 10
ABS29927/c
ID ABS29927 standard; DNA; 425 BP.
XX
AC ABS29927;
XX
DT 25-FEB-2003 (first entry)
XX
DE Human liver single exon probe, SEQ ID No 4917.
XX
KW Human; single exon nucleic acid probe; liver; cirrhosis;
KW hyperlipoproteinaemia; hyperlipidaemia; hypercholesterolaemia;
KW coronary heart disease; ss.
XX
OS Homo sapiens.
XX
PN WO200157273-A2.
XX
PD 09-AUG-2001.
XX
PF 30-JAN-2001; 2001WO-US00664.
XX
PR 04-FEB-2000; 2000US-0180312.
PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX Penn SG, Hanzel DK, Chen W, Rank DR;
XX WPI; 2001-488898/53.
XX Human genome-derived single exon nucleic acid probes useful for
PT analysing gene expression in human adult liver -
XX
PS Claim 1; SEQ ID No 4917; 658pp; English.
XX
XX The invention relates to a single exon nucleic acid probe (SENP) (I) for
CC measuring human gene expression in a sample derived from human adult
CC liver, comprising one of 13109 defined nucleotide sequences given in the
CC specification (or complements/ fragments). The probe hybridises at high
CC stringency to a nucleic acid molecule expressed in the human adult

CC liver. (1) may be used for predicting, measuring and displaying gene
 CC expression in samples derived from human adult liver. The genes
 CC identified may be involved in genetic liver diseases such as cirrhosis,
 CC hyperlipoproteinaemia, hyperlipidaemia and hypercholesterolaemia which
 CC is associated with coronary heart disease. AB25011-AB551005 represent
 CC human liver single exon nucleic acid probes of the invention.
 CC Note: The sequence information for this patent does not appear in the
 CC printed specification but was obtained in electronic format directly
 CC from WIPO at ftp.wipo.int/pub/published_pct_sequences.
 XX
 SQ Sequence 425 BP; 112 A; 107 C; 94 G; 112 T; 0 other;
 Query Match 100.0%; Score 21; DB 23; Length 425;
 Best Local Similarity 100.0%; Pred. No. 0.69;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 AAGCCTGATTGATGACGAGG 21
 DB 150 AAGCCTGATTGATGACGAGG 130
 RESULT 11
 ID ABS04866/c
 AC ABS04866 standard; DNA; 425 BP.
 AC ABS04866;
 XX
 DT 19-AUG-2002 (first entry)
 XX
 DE Human genome-derived single exon probe from lung SEQ ID No 4857.
 XX
 KW Human; ds; single exon probe; asthma; lung cancer; COPD; ILD;
 KW chronic obstructive pulmonary disease; interstitial lung disease;
 KW familial idiopathic pulmonary fibrosis; neurofibromatosis;
 KW tuberous sclerosis; Gaucher's disease; Niemann-Pick disease;
 KW Hermansky-Pudlak syndrome; sarcoidosis; pulmonary haemosiderosis;
 KW pulmonary histiocytosis; lymphangioleiomyomatosis; Karagener syndrome;
 KW pulmonary alveolar proteinosis; fibrocystic pulmonary dysplasia;
 KW primary ciliary dyskinesia; pulmonary hypertension;
 KW hyaline membrane disease.
 XX
 OS Homo sapiens.
 XX
 PN WO200186003-A2.
 XX
 PD 15-NOV-2001.
 XX
 PF 30-JAN-2001; 2001WO-US00665.
 XX
 PR 04-FEB-2000; 2000US-180312P.
 PR 26-MAY-2000; 2000US-207456P.
 PR 30-JUN-2000; 2000US-0608408.
 PR 03-AUG-2000; 2000US-0623366.
 PR 21-SEP-2000; 2000US-234687P.
 PR 27-SEP-2000; 2000US-236359P.
 PR 04-OCT-2000; 2000US-0024263.
 XX
 PA (MOLE-) MOLECULAR DYNAMICS INC.
 XX
 PI Penn SG, Hanzel DK, Chen W, Rank DR;
 XX
 XX WPI; 2002-114193/15.
 XX
 PT Spatially-addressable set of single exon nucleic acid probes, used to
 PT measure gene expression in human lung samples -
 XX
 XX Claim 1; SEQ ID No 4857; 634pp; English.
 XX
 CC The invention relates to a spatially-addressable set of single exon
 CC nucleic acid probes for measuring gene expression in a sample derived
 CC from human lung comprising single exon nucleic acid probes having one of
 CC 12614 nucleic acid sequences mentioned in the specification, or their
 CC complements or the 12387 open reading frames derived from the 12614

CC probes. Also included are a microarray comprising the novel set of
 CC probes; the novel set of probes which hybridise at high stringency to a
 CC nucleic acid expressed in the human lung; measuring gene expression in a
 CC sample derived from human lung, comprising (a) contacting the array with
 CC a collection of detectably labeled nucleic acids derived from human lung
 CC mRNA, and (b) measuring the label detectably bound to each probe of
 CC the array; identifying exons in a eukaryotic genome, comprising
 CC (a) algorithmically predicting at least one exon from genomic sequences
 CC of the eukaryote; and (b) detecting specific hybridisation of detectably
 CC labeled nucleic acids from eukaryote lung mRNA, to a single exon probe,
 CC having a fragment identical to the predicted exon, the probe is included
 CC in the above mentioned microarray; assigning exons to a single gene,
 CC comprising (a) identifying exons from genomic sequence by the method
 CC above and (b) measuring the expression of each of the exons in several
 CC tissues and/or cell types using hybridisation to a single exon
 CC microarrays having a probe with the exon, where a common pattern of
 CC expression of the exons in the tissues and/or cell types indicates that
 CC the exons should be assigned to a single gene; a peptide comprising one
 CC of 12011 sequences, mentioned in the specification, or encoded by the
 CC probes/open reading frames (ORF). The probes are used for gene
 CC expression analysis, and for identifying exons in a gene, particularly
 CC using human lung derived mRNA and for the study of lung diseases
 CC such as asthma, lung cancer, chronic obstructive pulmonary disease
 CC (COPD), interstitial lung disease (ILD), familial idiopathic pulmonary
 CC fibrosis, neurofibromatosis, tuberous sclerosis, Gaucher's disease,
 CC Niemann-Pick disease, Hermansky-Pudlak syndrome, sarcoidosis, pulmonary
 CC haemosiderosis, pulmonary histiocytosis, lymphangioleiomyomatosis,
 CC pulmonary alveolar proteinosis, Karagener syndrome, fibrocystic,
 CC pulmonary dysplasia, primary ciliary dyskinesia, pulmonary hypertension
 CC and hyaline membrane disease. The present sequence is a single exon
 CC probe of the invention.
 CC Note: The sequence data for this patent did not form part
 CC of the printed specification, but was obtained in electronic
 CC format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.
 XX
 SQ Sequence 425 BP; 112 A; 107 C; 94 G; 112 T; 0 other;
 Query Match 100.0%; Score 21; DB 24; Length 425;
 Best Local Similarity 100.0%; Pred. No. 0.69;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 AAGCCTGATTGATGACGAGG 21
 DB 150 AAGCCTGATTGATGACGAGG 130
 RESULT 12
 ABX95006
 ID ABX95006 standard; cDNA; 1041 BP.
 XX
 AC ABX95006;
 XX
 DT 05-JUN-2003 (first entry)
 XX
 DE cDNA encoding human tumour rejection antigen precursor, MAGE-C3.
 XX
 KW TRAP; ss; tumour rejection antigen precursor; cytolytic T-cell; CTL;
 KW tumour; seminoma; bladder transitional-cell carcinoma; NSCLC; adaptor;
 KW head-and-neck squamous-cell carcinoma; breast carcinoma; sarcoma;
 KW cutaneous melanoma; non-small cell lung cancer; gene; MAGE-C3; human;
 KW chromosome Xq27.1-Xq27.3.
 XX
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 CDS 1..1041
 FT /*tag= a
 FT /product= "MAGE-C3"
 XX
 PN US2002176865-A1.
 XX
 PD 28-NOV-2002.

XX PF 01-MAR-2002; 2002US-0085108.
 XX XX
 XX XX
 PR 09-FEB-2000; 2000US-0501104.
 PR 25-APR-1997; 97US-0845528.
 PR 24-APR-1998; 98US-0066281.
 PR 17-DEC-1999; 99US-0468433.
 XX XX
 PA (LUCAS/) LUCAS S.
 PA (BOON/) BOON-FALLEUR T.
 XX XX
 XX Lucas S, Boon-Falleur T;
 PI XX
 XX WPI: 2003-328468/31.
 DR P-PSDB; ABU08932.
 DR XX
 XX Novel isolated nucleic acid encoding tumor rejection antigen precursor
 PT MAGE-C3, MAGE-B5, or MAGE-B6, useful as diagnostic probes to determine
 PT presence of abnormal e.g., tumor cells expressing MAGE-C1, MAGE-B5 or
 PT MAGE-B6 -
 XX XX
 PS Claim 1; Fig 5; 59pp; English.
 XX
 CC The invention relates to an isolated nucleic acid molecule which encodes
 CC a tumour rejection antigen precursor (TRAP) having an amino acid sequence
 CC of a TRAP encoded by a fully defined MAGE-C3, MAGE-B5, or MAGE-B6
 CC polynucleotide sequence. Also disclosed is a method which is useful for
 CC determining presence of cytolytic T-cells specific for complexes of human
 CC leukocyte antigen (HLA) and a peptide derived from the nucleic acid in a
 CC cytotoxic T-lymphocyte (CTL)-containing sample. The nucleic acid is
 CC useful as a diagnostic probe to determine the presence of abnormal
 CC (tumour) cells such as seminoma, bladder transitional-cell carcinoma,
 CC head-and-neck squamous-cell carcinoma, breast carcinoma, sarcoma,
 CC cutaneous melanoma or non-small cell lung cancer (NSCLC) which express
 CC MAGE-C1, MAGE-B5 or MAGE-B6. The nucleic acid is useful for diagnosing a
 CC disorder characterised by expression of MAGE-C1, MAGE-B5 or MAGE-B6 TRAPS
 CC or tumour rejection antigens (TRAs). The present sequence represents the
 CC cDNA of the gene encoding the human tumour rejection antigen precursor,
 CC MAGE-C3, which is located on chromosome Xq27.1-Xq27.3.
 XX
 SQ Sequence 1041 BP; 242 A; 283 C; 242 G; 274 T; 0 other;
 Query Match 100.0%; Score 21; DB 25; Length 1041;
 Best Local Similarity 100.0%; Pred. No. 0.81; Mismatches 0; Indels 0; Gaps 0;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 AAGCCTGATTGATGACCAGG 21
 DB 711 AAGCCTGATTGATGACCAGG 731
 RESULT 13
 ID AAS88354
 XX AAS88354 standard; cDNA; 7806 BP.
 AC AAS88354;
 XX
 DT 13-FEB-2002 (first entry)
 XX
 DE DNA encoding novel human diagnostic protein #24158.
 XX
 KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200175067-A2.
 XX
 PD 11-OCT-2001.
 XX
 PF 30-MAR-2001; 2001WO-US08631.
 XX
 PR 31-MAR-2000; 2000US-0540217.

PR 23-AUG-2000; 2000US-0649167.
 XX (HYSE-) HYSEQ INC.
 XX
 XX Drmanac RT, Liu C, Tang YT;
 XX
 DR WPI: 2001-639362/73.
 DR P-PSDB; ABG24167.
 XX
 XX New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -
 XX
 PS Claim 1; SEQ ID No 24158; 103pp; English.
 XX
 CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 XX
 SQ Sequence 7806 BP; 1755 A; 2185 C; 1760 G; 2106 T; 0 other;
 Query Match 100.0%; Score 21; DB 23; Length 7806;
 Best Local Similarity 100.0%; Pred. No. 1.2; Mismatches 0; Indels 0; Gaps 0;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 AAGCCTGATTGATGACCAGG 21
 DB 3971 AAGCCTGATTGATGACCAGG 3991
 RESULT 14
 AAT24776
 ID AAT24776 standard; cDNA to mRNA; 336 BP.
 XX
 XX AAT24776;
 AC AAT24776;
 XX
 DT 03-OCT-1996 (first entry)
 XX
 DE Human gene signature HUMGS06854.
 XX
 KW Gene signature; messenger RNA; mRNA; relative abundance; frequency;
 KW human; cloning; mapping; non-biased library; diagnosis; detection;
 KW cell typing; abnormal cell function; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO9514772-A1.
 XX
 PD 01-JUN-1995.
 XX
 PF 11-NOV-1994; 94WO-JP01916.
 XX
 PR 12-NOV-1993; 93JP-0355504.
 XX

PA (MATS/) MATSUBARA K.
 XX (OKUB/) OKUBO K.
 XX Matsubara K, Okubo K;
 PI WPI; 1995-206931/27.
 XX
 DR Identifying gene signatures in 3'-directed human cDNA library - e.g.
 PT for diagnosis of abnormal cell function, by preparing cDNA that
 PT reflects relative abundance of corresp. mRNA in specific human
 PT tissues
 XX
 PS Claim 1; Page 1692; 2245pp; Japanese.
 XX
 CC A single-stranded DNA (or its complementary strand or the corresp.
 CC double-stranded DNA) which comprises one of the 7837 "GS" sequences
 CC given in AAT19001-T26837 and which is able to hybridise to part of
 CC human genomic DNA, cDNA or mRNA is claimed. The GS (Gene Signature)
 CC sequences were obtained from 3'-directed cDNA libraries prepared
 CC from various human tissues; synthesis of cDNA was initiated from the
 CC 3'-end of mRNA by using poly(T) as the sole primer. Since the 3'-
 CC untranslated sequence is unique to a particular mRNA species, almost
 CC all the 3'-oriented cDNAs hybridise with specific mRNAs. Each library
 CC is constructed so as to reflect accurately the relative abundance of
 CC different mRNAs in the particular tissue from which it was derived.
 CC The appearance frequency of a given GS in a cDNA library can be
 CC determined (esp. using primers and probes derived from the GS
 CC sequences) as a means of diagnosing abnormal cell function or for
 CC recognising different cell types.
 XX
 SQ Sequence 336 BP; 91 A; 67 C; 80 G; 83 T; 15 other;
 Query Match 81.0%; Score 17; DB 16; Length 336;
 Best Local Similarity 100.0%; Pred. No. 73;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 5 CTGATTGATGACCGGG 21
 Db 231 CTGATTGATGACCGGG 247
 RESULT 15
 ABS69805
 ID ABS69805 standard; DNA; 504 BP.
 XX
 AC ABS69805;
 XX
 DT 21-NOV-2002 (first entry)
 XX
 DE Novel murine polynucleotide isolated using gene trap technology #869.
 XX
 KW Mouse; gene trapped sequence; GTS; functional genomic analysis;
 KW phage display system; gene chip; temporal gene expression;
 KW tissue specific gene expression; antisense inhibition; gene targeting;
 KW development disorder; cell differentiation disorder; aging; cancer;
 KW autoimmune disease; lupus; inflammatory disorder; skin disorder;
 KW degenerative disorder; de.
 XX
 OS Mus musculus.
 XX
 XX US2002102543-A1.
 PN
 XX
 PD 01-AUG-2002.
 XX
 XX 30-NOV-2000; 2000US-0728445.
 PF
 XX
 PR 01-DEC-1999; 99US-168358P.
 XX
 XX (FRIE/) FRIEDRICH G.
 PA (ZAMB/) ZAMBROWICZ B.
 PA (SAND/) SANDS A T.
 XX
 PI Friedrich G, Zambrowicz B, Sands AT;
 WPI; 2002-690598/74.
 DR Novel murine polynucleotides that individually identify novel genes
 PT into which a retroviral gene trap vector has integrated, useful in
 PT genomic analysis and in discovery, development of therapeutic and
 PT diagnostic agents -
 XX
 PS Claim 1; Page 286; 296pp; English.
 XX
 CC The invention describes an isolated murine polynucleotide (I) comprising
 CC a contiguous stretch of at least 60 nucleotides of one of 265-677
 CC nucleotide 891 OMNIBANK gene trapped sequences (GTSs) (S), given in the
 CC specification. The novel genes and cells are useful in functional
 CC genomic analysis and in the discovery and development of new therapeutic
 CC and diagnostic agents and methods. (I) is useful for identifying the
 CC coding regions of the murine genome, to isolate cDNAs, genomic clones,
 CC or full-length genes/polynucleotides or homologues, heterologues,
 CC paralogues, or orthologues that are capable of hybridising to one or more
 CC of the GTSs under stringent conditions. (I) can be incorporated into a
 CC phage display system that can be used to screen for proteins or other
 CC ligands, that are capable of binding an amino acid sequence encoded by
 CC an oligonucleotide or polynucleotide sequence in at least one of the TS
 CC sequences. (I) is useful in addressable arrays, such as gene chips, to
 CC identify and characterise temporal and tissue specific gene expression,
 CC to identify the gene of interest from many sources and for genetic
 CC manipulations such as antisense inhibition and gene targeting. Decreasing
 CC the level of expression of (I) and/or down regulating the activity of
 CC peptides or proteins encoded by (I) is useful for treating development
 CC and cell differentiation disorders, aging, cancer, autoimmune disease,
 CC lupus, inflammatory disorders, skin disorders and degenerative
 CC disorders. This sequence represents a murine cDNA isolated using gene
 CC trap technology.
 XX
 SQ Sequence 504 BP; 110 A; 152 C; 146 G; 96 T; 0 other;
 Query Match 80.0%; Score 16.8; DB 24; Length 504;
 Best Local Similarity 90.0%; Pred. No. 1e+02;
 Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 2 AGCCTGATTGATGACCGGG 21
 Db 461 AGCAGGATTGATGACCGGG 480
 Search completed: February 19, 2004, 21:28:33
 Job time : 176.5 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: February 19, 2004, 20:01:08 ; Search time 1596.5 Seconds
(without alignments)
538.116 Million cell updates/sec

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Perfect score: 21
Sequence: 1 AAGCCTGATTGATGACCAGGG 21

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 2888711 seqs, 2045481386 residues

Total number of hits satisfying chosen parameters: 5777422

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : GenEmbl.*

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- 2: gb.htg.*
- 3: gb.in.*
- 4: gb.om.*
- 5: gb.ov.*
- 6: gb.pat.*
- 7: gb.ph.*
- 8: gb.pl.*
- 9: gb.pr.*
- 10: gb.ro.*
- 11: gb.sts.*
- 12: gb.sy.*
- 13: gb.un.*
- 14: gb.vi.*
- 15: em.ba.*
- 16: em.fun.*
- 17: em.hum.*
- 18: em.in.*
- 19: em.mu.*
- 20: em.om.*
- 21: em.or.*
- 22: em.ov.*
- 23: em.pat.*
- 24: em.ph.*
- 25: em.pl.*
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- 27: em.sts.*
- 28: em.un.*
- 29: em.vi.*
- 30: em.htg.hum.*
- 31: em.htg.inv.*
- 32: em.htg.other.*
- 33: em.htg.mus.*
- 34: em.htg.pln.*
- 35: em.htg.rod.*
- 36: em.htg.man.*
- 37: em.htg.vrt.*
- 38: em.sy.*
- 39: em.htgo.hum.*
- 40: em.htgo.mus.*
- 41: em.htgo.other.*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB	ID	Description
1	21	100.0	1932	9	AF490508	AF490508 Homo sapi
2	21	100.0	124497	9	HS232G24	AL022152 Human DNA
C 3	19	90.5	153439	2	EX323047	BX323047 Danio rer
C 4	19	90.5	159948	2	BX248240	BX248240 Danio rer
5	17.8	84.8	1494	9	AK098443	AK098443 Homo sapi
6	17.8	84.8	1681	9	BC048268	BC048268 Homo sapi
C 7	17.8	84.8	67201	9	AL136169	AL136169 Human DNA
8	17.8	84.8	163746	2	AC102710	AC102710 Mus muscu
C 9	17.8	84.8	174191	9	AC109462	AC109462 Homo sapi
C 10	17.8	84.8	177733	2	AC113908	AC113908 Rattus no
C 11	17.8	84.8	180353	2	AC128288	AC128288 Rattus no
C 12	17.8	84.8	186118	2	AC117856	AC117856 Rattus no
C 13	17.8	84.8	194454	2	AC111040	AC111040 Mus muscu
C 14	17.8	84.8	194520	10	AL691424	AL691424 Mouse DNA
15	17.8	84.8	194636	10	AC123058	AC123058 Mus muscu
C 16	17.8	84.8	214493	2	AC097047	AC097047 Rattus no
C 17	17.8	84.8	227369	2	AC114469	AC114469 Rattus no
C 18	17.8	84.8	231827	2	AC096130	AC096130 Rattus no
C 19	17.8	84.8	232476	2	AC18621	AC18621 Mus muscu
C 20	17.8	84.8	243257	2	AC130512	AC130512 Rattus no
21	17.8	84.8	251759	2	AC137283	AC137283 Rattus no
C 22	17.8	84.8	257104	2	AC122547	AC122547 Mus muscu
C 23	17.8	84.8	274024	2	AC102995	AC102995 Rattus no
C 24	17.8	84.8	274467	2	AC113707	AC113707 Rattus no
C 25	17.4	82.9	196	5	AY185781	AY185781 Ambystoma
C 26	17.4	82.9	197	5	AY185782	AY185782 Ambystoma
C 27	17.4	82.9	223	5	AY185785	AY185785 Ambystoma
C 28	17.4	82.9	223	5	AY185786	AY185786 Ambystoma
C 29	17.4	82.9	224	5	AY185783	AY185783 Ambystoma
C 30	17.4	82.9	224	5	AY185787	AY185787 Ambystoma
C 31	17.4	82.9	227	5	AY185784	AY185784 Ambystoma
C 32	17.4	82.9	4202	5	AF001958	AF001958 Ambystoma
C 33	17.4	82.9	92337	10	AL604022	AL604022 Mouse DNA
C 34	17.4	82.9	120492	3	CEY43F8C	AL032637 Caenorhab
C 35	17.4	82.9	198727	2	AC122401	AC122401 Mus muscu
C 36	17.4	82.9	203651	9	AC011374	AC011374 Homo sapi
C 37	17.4	82.9	218092	2	AC019272	AC019272 Mus muscu
C 38	17.4	82.9	219451	2	AC137244	AC137244 Rattus no
39	17.4	82.9	226771	2	AC110940	AC110940 Rattus no
40	17.4	82.9	242602	2	AC095427	AC095427 Rattus no
41	17	81.0	108542	10	AL627307	AL627307 Mouse DNA
42	17	81.0	208966	2	AC099588	AC099588 Mus muscu
C 43	17	81.0	212119	2	AC099602	AC099602 Mus muscu
C 44	17	81.0	236604	2	AC099619	AC099619 Mus muscu
C 45	16.8	80.0	403	9	AF490961	AF490961 Homo sapi

ALIGNMENTS

RESULT 1
AF490508
LOCUS
DEFINITION Homo sapiens hepatocellular carcinoma-associated protein HCA2 mRNA,
complete cds.
ACCESSION AF490508
VERSION AF490508.1 GI:19919741
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 1932)
AUTHORS Dong,X. and Chen,W.
TITLE Identification of genes in the chromosome X that are differentially

 * NOTE: This is a 'working draft' sequence. It currently
 * consists of 13 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

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1      3298: contig of 3298 bp in length
*      3299: gap of 100 bp
*      3398: contig of 7210 bp in length
*      3399: contig of 100 bp
*      10609: gap of 27115 bp in length
*      37823: gap of 100 bp
*      37924: contig of 7452 bp in length
*      45375: gap of 100 bp
*      45476: gap of 100 bp
*      58730: contig of 13255 bp in length
*      58731: gap of 100 bp
*      58830: gap of 100 bp
*      61226: contig of 2396 bp in length
*      61327: gap of 100 bp
*      61327: gap of 100 bp
*      64481: contig of 3155 bp in length
*      64482: gap of 100 bp
*      84036: contig of 19455 bp in length
*      84037: gap of 100 bp
*      84136: gap of 100 bp
*      89161: contig of 5025 bp in length
*      89162: gap of 100 bp
*      89262: contig of 24645 bp in length
*      113906: contig of 100 bp
*      114007: gap of 100 bp
*      114007: contig of 28013 bp in length
*      142019: gap of 100 bp
*      142020: gap of 100 bp
*      149275: contig of 7156 bp in length
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*      149376: 153439: contig of 4064 bp in length.

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FEATURES

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37924..45375
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61327..64481
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89262..113906
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BASE COUNT

Query Match

90.5%; Score 19; DB 2; Length 153439;

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Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 3 GCCTGATTGATGACACAGG 21
Db 138696 GCCTGATTGATGACACAGG 138678

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RESULT 4

BX248240/c

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LOCUS      BX248240      159948 bp      DNA      linear      HTG 03-APR-2003
DEFINITION Danio rerio clone CH211-162119, *** SEQUENCING IN PROGRESS ***, 8
            unordered pieces.
ACCESSION  BX248240.3  GI:29539163
VERSION    BX248240.3  GI:29539163
KEYWORDS   HTG; HTGS PHASE1; HTGS DRAFT; HTGS_FULLTOP.
SOURCE     Danio rerio (zebrafish)
ORGANISM   Danio rerio
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
            Cypriniformes; Cyprinidae; Danio.
REFERENCE  1 (bases 1 to 159948)
AUTHORS    McLaren,S.
TITLE      Direct Submission
JOURNAL    Submitted (03-APR-2003) Wellcome Trust Sanger Institute, Hinxton,
            Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
            zfish-help@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
            On Apr 3, 2003 this sequence version replaced gi:28268576.
COMMENT    ----- Genome Center
            Center: Wellcome Trust Sanger Institute
            Center code: SC
            Web site: http://www.sanger.ac.uk
            Contact: zfish-help@sanger.ac.uk
            ----- Project Information
            Center project name: ZC162119
            ----- Summary Statistics
            Assembly program: XGAP4; version 4.5
            Chemistry: Dye-terminator; 100% of reads
            Consensus quality: 156495 bases at least Q40
            Consensus quality: 157186 bases at least Q30
            Consensus quality: 158113 bases at least Q20
            Insert size: 159248; sum-of-contigs
            Insert size: 152025; 6.7% error; agarose-fp
            Quality coverage: 9.26x in Q20 bases; sum-of-contigs Quality
            coverage: 9.80x in Q20 bases; agarose-fp
            -----
            * NOTE: This is a 'working draft' sequence. It currently
            * consists of 8 contigs. The true order of the pieces
            * is not known and their order in this sequence record is
            * arbitrary. Gaps between the contigs are represented as
            * runs of N, but the exact sizes of the gaps are unknown.
            * This record will be updated with the finished sequence
            * as soon as it is available and the accession number will
            * be preserved.
            1      3108: contig of 3108 bp in length
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            *      3208: gap of 100 bp
            *      32890: contig of 29682 bp in length
            *      32891: gap of 100 bp
            *      32990: gap of 100 bp
            *      32991: gap of 100 bp
            *      49822: contig of 16832 bp in length
            *      49823: gap of 100 bp
            *      49923: contig of 5141 bp in length
            *      55064: gap of 100 bp
            *      55164: contig of 56681 bp in length
            *      111844: gap of 100 bp
            *      111845: gap of 100 bp
            *      131419: contig of 19475 bp in length
            *      131420: gap of 100 bp
            *      131519: gap of 100 bp
            *      149030: contig of 17511 bp in length
            *      149031: gap of 100 bp
            *      149130: gap of 100 bp
            *      149131: 159948: contig of 10818 bp in length.
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            /mol_type="genomic DNA"

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FEATURES

source

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1..159948
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32991 .49822
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  fragment_chain:1"
49923 .55063
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  fragment_chain:1"
55164 .111844
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  fragment_chain:1"
111945 .131419
  /note="assembly_fragment:00155
  fragment_chain:1"
131520 .149030
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ORIGIN

Query Match 90.5%; Score 19; DB 2; Length 159948;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AAGCTGATTGATGACCAG 19
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Db 92696 AAGCTGATTGATGACCAG 92678
|||||

RESULT 5
AK098443 1494 bp mRNA linear PRI 15-JUL-2002
LOCUS
DEFINITION Homo sapiens cDNA FLJ25577 fis, clone JTH07710.
ACCESSION AK098443
VERSION AK098443 1 GI:21758456
KEYWORDS Oligo capping; fis (full insert sequence).
SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE
1
AUTHORS Ninomiya,K., Wagatsuma,M., Kanda,K., Kondo,H., Yokoi,T.,
Kodaira,H., Furiya,T., Takahashi,M., Kikkawa,B., Omura,Y., Abe,K.,
Kaminara,K., Katsuta,N., Sato,K., Tanikawa,M., Yamazaki,M.,
Suzuki,Y., Hata,H., Nakagawa,K., Mizuno,S., Morinaga,M.,
Kawamura,M., Sugiyama,T., Irie,R., Otsuki,T., Sato,H.,
Nishikawa,T., Sugiyama,A., Kawakami,B., Negai,K., Isogai,T. and
Sugano,S.
NEDO human cDNA sequencing project
TITLE NEDO human cDNA sequencing project
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 1494)
AUTHORS Sugano,S. and Suzuki,Y.
TITLE Direct Submission
JOURNAL Submitted (08-JUL-2002) Sumio Sugano, Institute of Medical Science,
University of Tokyo, Laboratory of Genome Structure, Human Genome
Center, Shirokane-dai, 4-6-1, Minato-ku, Tokyo 108-8639, Japan
(E-mail:cdna@ims.u-tokyo.ac.jp, Tel:81-3-5449-5286,
Fax:81-3-5449-5416)
COMMENT NEDO human cDNA sequencing project supported by Ministry of
Economy, Trade and Industry of Japan; cDNA full insert sequencing:
Research Association for Biotechnology (RAB); cDNA library

construction and 5'-end one pass sequencing: Institute of Medical
Science, University of Tokyo, Laboratory of Genome Structure, Human
Genome Center; 3'-end one pass sequencing: RAB; clone selection for
full insert sequencing: RAB and Helix Research Institute.
FEATURES
source
Location/Qualifiers
1 .1494
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="JTH07710"
/cell_line="JCR"
/tissue_type="thyroid"
/clone_lib="JTH"
/note="cloning vector: pME18SPL3"
BASE COUNT 368 a 394 c 292 g 440 t
ORIGIN

Query Match 84.8%; Score 17.8; DB 9; Length 1494;
Best Local Similarity 90.5%; Pred. No. 5.5e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 AAGCTGATTGATGACCAGG 21
|||||
Db 1244 AACCTGGTTGATGACCAGG 1264
|||||

RESULT 6
BC048268 1681 bp mRNA linear PRI 31-MAR-2003
LOCUS
DEFINITION Homo sapiens, clone IMAGE:4512974, mRNA.
ACCESSION BC048268
VERSION BC048268.1 GI:29387246
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE
1 (bases 1 to 1681)
AUTHORS Strausberg,R.
TITLE Direct Submission
JOURNAL Submitted (07-MAR-2003) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics Office, National Cancer
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
USA
NTH-MGC project URL: http://mgc.nci.nih.gov
Contact: MGC help desk
Email: gcgbs-remail.nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: Life Technologies, Inc.
DNA Library Arrayed by: The I.M.A.G.E. Consortium (ILLNL)
DNA Sequencing by: National Institutes of Health Intramural
Sequencing Center (NISC),
Gaithersburg, Maryland;
Web site: http://www.nisc.nih.gov/
Contact: nisc.mgc@nih.gov
Contact: nisc.mgc@nih.gov
Blakesley,R.W., Bouffard,G.G., Breen,K., Brinkley,C., Brooks,S.,
Dietrich,N.L., Granite,S., Guan,X., Gupta,J., Haghighi,P.,
Hansen,N., Ho,S.-L., Karlins,E., Kwong,P., Laric,P., Legaspi,R.,
Maduro,Q.L., Masiello,C., Maskeri,B., Mastrian,S.D., McCloskey,J.C.,
McDowell,J., Pearson,R., Stantropop,S., Thomas,P.J., Touchman,J.W.,
Teurgeon,C., Vogt,J.L., Walker,M.A., Wetherby,K.D., Wiggins,L.,
Young,A., Zhang,L.-H. and Green,E.D.
Clone distribution: MGC clone distribution information can be found
through the I.M.A.G.E. Consortium/ILLNL at: http://image.llnl.gov
Series: IRAC Plate: 93 Row: d Column: 5
This clone was selected for full length sequencing because it
passed the following selection criteria: Similarity but not
identity to protein.
FEATURES
source
Location/Qualifiers
1 .1681
/organism="Homo sapiens"

```

```

/molecule="mrna"
/db_xref="taxon:9606"
/clone="IMAGE:4512974"
/tissue_type="Bladder, transitional cell papilloma"
/clone_lib="NIH_MGC_93"
/lab_host="DH10B"
/notes="Vector: pCMV-SPORT6"

BASE COUNT      404 a      449 c      357 g      471 t
ORIGIN
Query Match      84.8%; Score 17.8; DB 9; Length 1681;
Best Local Similarity 90.5%; Pred. No. 5.5e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      1 AAGCTGATTCATGACACGAGG 21
      |||||
Db      1414 AAACCTGGTTCATGACACGAGG 1434

RESULT 7
LOCUS      AL136169          67201 bp      DNA      linear      PRI 05-JUN-2003
DEFINITION Human DNA sequence from clone RP4-809E13 on chromosome Xq26.1-27.1
Contains the gene for a putative prenylated protein, two putative
prenyated protein pseudogenes, ESTs, STSS, GSSs and three putative
CpG islands, complete sequence.
ACCESSION      AL136169
VERSION      AL136169.6 GI:1760488
KEYWORDS      HTG; CpG island.
SOURCE      Homo sapiens (human)
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Heath,P.
REFERENCE      Direct Submission
Submitted (05-JUN-2003) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
humquerry@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
On Mar 6, 2000 this sequence version replaced GI:6996160.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases:
EMBL, EMBL; SW, SWISSPROT; Tr, TREMBL; Wp, WORMPEP; Information
on the WORMPEP database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence
was generated from part of bacterial clone contigs of human
chromosome X, constructed by the Sanger Centre Chromosome X Mapping
Group. Further information can be found at
http://www.sanger.ac.uk/HGP/ChrX
----- Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: humquerry@sanger.ac.uk
-----
This sequence was finished as follows unless otherwise noted: all
regions were either double-stranded or sequenced with an alternate
chemistry or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by at least
one plasmid subclone or more than one M13 subclone; and the
assembly was confirmed by restriction digest, except on the rare
occasion of the clone being a YAC.
RP4-809E13 is from the library RPCI-4 constructed by the group of
Pieter de Jong. For further details see
http://www.chori.org/bacpac/home.htm
VECTOR: pCYPAC2

```

IMPORTANT: This sequence is not the entire insert of clone RP4-809E13. It may be shorter because we sequence overlapping sections only once, except for a short overlap. The true right end of clone RP4-809E13 is at 67201 in this sequence. The true right end of clone RP3-473B4 is at 100 in this sequence.

FEATURES	Location/Qualifiers
source	1..67201
	/organism="Homo sapiens"
	/mol_type="genomic DNA"
	/db_xref="taxon:9606"
	/chromosome="X"
	/map="q26.1-27.1"
	/clone="RP4-809E13"
	/clone_lib="RPCI-4"
misc_feature	1..159
	/note="match: GSS: Em:B42366"
	/complement(1..74)
misc_feature	550..597
	/note="match: STS: Em:AL008947 Em:HS473B4T"
repeat_region	550..597
	/note="2 copies 24 mer 95% conserved"
repeat_region	758..922
	/note="LIME repeat: matches 5531..5695 of consensus"
repeat_region	939..1213
	/note="L2 repeat: matches 1806..2096 of consensus"
repeat_region	1271..1565
	/note="AluX repeat: matches 1..295 of consensus"
repeat_region	2011..2163
	/note="LIM4 repeat: matches 3743..3917 of consensus"
repeat_region	2642..2689
	/note="2 copies 24 mer 97% conserved"
repeat_region	3206..3274
	/note="3 copies 23 mer 81% conserved"
repeat_region	3243..3418
	/note="88 copies 2 mer aa 56% conserved"
repeat_region	3534..3636
	/note="MIR repeat: matches 16..115 of consensus"
repeat_region	4253..4518
	/note="AluJo repeat: matches 9..273 of consensus"
repeat_region	5225..5524
	/note="AluX repeat: matches 1..299 of consensus"
repeat_region	7872..8066
	/note="MBR67C repeat: matches 1..200 of consensus"
misc_feature	complement(11087..11278)
	/note="match: GSS: Em:AQ283138"
repeat_region	14329..14451
	/note="FAMC repeat: matches 1..126 of consensus"
repeat_region	15570..15671
	/note="MIR repeat: matches 110..218 of consensus"
misc_feature	17271..17692
	/note="match: GSS: Em:AQ182699"
misc_feature	17275..17984
	/note="match: GSS: Em:AQ553839"
repeat_region	18475..18708
	/note="LIPAS repeat: matches 5912..6141 of consensus"
repeat_region	18711..18752
	/note="21 copies 2 mer aa 78% conserved"
misc_feature	18965..19171
	/note="match: STS: Em:AA894450"
repeat_region	19099..19303
	/note="LTR37B repeat: matches 256..464 of consensus"
gene	complement(19859..20487)
	/genes="dJ809E13.1"
	/pseudo
CDS	complement(19859..20487)
	/genes="dJ809E13.1"
	/note="dJ809E13.1 [putative prenylated protein pseudogene]"
	match: cDNAs: Em:AB019527 Em:AF086537 Em:L76224 Em:U77782
	match: ESTs: Em:AI815482 Em:AI573275 Em:AA659153 Em:N72726
	Em:AI183679 Em:AW370707 Em:AW370768
	match: proteins: SW:P20824 Tr:O88409 Sw:P11675 Tr:O95751
	Tr:Q15779 Tr:Q6882 Tr:Q70555 Tr:Q61824 Tr:Q63729
	Tr:Q9Y172 Tr:Q68872 Tr:O15255 Tr:Q14468 Tr:Q9YDA0

Sw:P33479 Sw:Q03391"
/pseudo
/evidence=not_experimental
20312..20917
/note="CpG island"
/evidence=not_experimental
21213..21450
/note="L1MC4 repeat: matches 7288..7525 of consensus"
21645..21701
/note="L1 repeat: matches 3880..3935 of consensus"
21704..21981
/note="MLT1D repeat: matches 260..568 of consensus"
21963..22196
/note="MLT1D repeat: matches 73..323 of consensus"
22197..23067
/note="MER11D repeat: matches 1..897 of consensus"
23068..23143
/note="MLT1D repeat: matches 1..73 of consensus"
23154..23260
/note="L1 repeat: matches 3759..3870 of consensus"
23261..23532
/note="L1MC2 repeat: matches 5757..6048 of consensus"
23510..23715
/note="L1MC/D repeat: matches 5411..5596 of consensus"
23816..24409
/note="L1MC/D repeat: matches 4796..5397 of consensus"
24410..24486
/note="MER2 repeat: matches 268..345 of consensus"
24496..24630
/note="L1 repeat: matches 4189..4323 of consensus"
24643..24784
/note="L1P repeat: matches 3712..3855 of consensus"
24774..25830
/note="L1PB2 repeat: matches 4333..5439 of consensus"
25828..25903
/note="L1P1 repeat: matches 5642..5717 of consensus"
25904..26345
/note="L1HS repeat: matches 5705..6146 of consensus"
26366..27087
/note="L1PB2 repeat: matches 5423..6155 of consensus"
27113..27322
/note="MER2 repeat: matches 1..212 of consensus"
27323..28033
/note="L1M4 repeat: matches 4063..4797 of consensus"
28034..28237
/note="L1PA12 repeat: matches 5944..6158 of consensus"
28238..28361
/note="L1M4 repeat: matches 3943..4063 of consensus"
28361..28536
/note="L1MC/D repeat: matches 5613..5792 of consensus"
28537..28689
/note="MLT1E repeat: matches 1..174 of consensus"
28690..29650
/note="LTP5 repeat: matches 1..969 of consensus"
29651..29765
/note="MLT1E repeat: matches 174..282 of consensus"
29848..30076
/note="L1M42 repeat: matches 6073..6226 of consensus"
30077..30137
/note="L1M41 repeat: matches 6169..6226 of consensus"
30383..31300
/note="CpG island"
/evidence=not_experimental
30658..31891
/gene="dJ809E13.2"
30658..31891
/gene="dJ809E13.2"
/product="dJ809E13.2 (putative prenylated protein)"
/note="match: cDNAs: Em:AF052096 Em:AF038168 Em:Y13374
match: ESTs: Em:AI742011 Em:F21403 Em:W76590 Em:AA307887
Em:F21483 Em:N64112 Em:AA421975 Em:AA031541 Em:AA122144
Em:AW410876 Em:AW370386 Em:F37666 Em:AI027000 Em:AA091701

Em:AA302562 Em:AA070647 Em:AA733171 Em:AA076516 Em:C04431
Em:AA864654 Em:AI339190 Em:AI086752 Em:AI346926

Query Match 84.8%; Score 17.8; DB 9; Length 67201;
Best Local Similarity 90.5%; Fred. No. 4.8e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAGCTGATTGATGACCAGGG 21
|||||
Db 19111 AAACCTGGTGTGACCAGGG 19091
|||||

RESULT 8
LOCUS AC102710
DEFINITION Mus musculus clone RP24-160E14, WORKING DRAFT SEQUENCE, 9 unordered
pieces.
AC102710
VERSION AC102710.3 GI:28395973
KEYWORDS HTGS PHASE1; HTGS DRAFT; HTGS_FULLTOP.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS
1 (bases 1 to 163746)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N.,
Anderson,S., Barna,N., Bastien,V., Boguslavsky,L., Bouckgalter,B.,
Brown,A., Camarata,J., Campopiano,A., Chang,J., Chazaro,B.,
Choepel,Y., Collangelo,M., Collins,S., Collymore,A., Cook,A.,
Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S.,
Ferreira,P., FitzHugh,W., Gage,D., Galagan,J., Gardyna,S.,
Gande,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N.,
Hagos,B., Heaford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R.,
Jones,C., Kamat,A., Karatas,A., Kells,C., LaRocque,K.,
Lamaze,R., Landers,T., Lehotzky,J., Levine,R., Liu,G.,
Maclean,C., Macdonald,P., Major,J., Marquis,N., Matthews,C.,
McCarthy,M., McEwan,P., McKernan,K., McPheters,R., Meldrim,J.,
Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C.,
Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D.,
Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V.,
Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P.,
Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S., Schupack,R.,
Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
Topham,K., Travers,M., Travis,N., Trigilio,J., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,
Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

Direct Submission
Submitted (23-NOV-2001) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 163746)
TITLE
JOURNAL
REFERENCE
AUTHORS
1 (bases 1 to 163746)
Birren,B., Nusbaum,C., Lander,E., Abouelleil,A., Allen,N.,
Anderson,S., Arachchi,H.M., Barna,N., Bastien,V., Bloom,T.,
Boguslavsky,L., Bouckgalter,B., Camarata,J., Chang,J., Choepel,Y.,
Collymore,A., Cooke,A., Cooke,P., Corum,B., Dearellano,K.,
Diaz,J.S., Dodge,S., Dooley,K., Dorris,L., Erickson,J., Faro,S.,
Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J., Gardyna,S.,
Graham,L., Grand-Pierre,N., Hafez,N., Hagopian,D., Hagos,B.,
Hall,J., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C.,
Kamat,A., Karatas,A., Kells,C., Landers,T., Levine,R.,
Lindblad-Toh,K., Liu,G., Lui,A., Mabbitt,R., Maclean,C.,
Macdonald,P., Major,J., Manning,J., Matthews,C., McCarthy,M.,
Meldrim,J., Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J.,
Nguyen,C., Nicol,R., Norbu,C., O'Connor,T., O'Donnell,P.,
O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N.,
Rachupka,A., Ramasamy,U., Raymond,C., Retta,R., Rise,C., Rogov,P.,
Roman,J., Schauer,S., Schupack,R., Seaman,S., Severy,P., Smith,C.,
Spencer,B., Stange-Thomann,N., Stojanovic,N., Stubbs,M.,
Talamas,J., Tesfaye,S., Theodore,J., Topham,K., Travers,M.,

TITLE
JOURNAL
COMMENT

Vassiliev, H., Venkataraman, V.S., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.
Submitted (17-FEB-2003) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
On Feb 17, 2003 this sequence version replaced gi:22381702.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: <http://www-seq.wi.mit.edu>

Contact: sequence_submissions@genome.wi.mit.edu

----- Project Information

Center project name: L19331

Center clone name: 160 E.14

----- Summary Statistics

Sequencing vector: Plasmid; n/a; 100% of reads

Chemistry: Dye-terminator Big Dye; 100% of reads

Assembly program: Phrap; version 0.960731

Consensus quality: 162090 bases at least Q40

Consensus quality: 162587 bases at least Q30

Consensus quality: 162779 bases at least Q20

Insert size: 157000; agarose-fp

Insert size: 162946; sum-of-contigs

Quality coverage: 9.4 in Q20 bases; agarose-fp

Quality coverage: 9.0 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 9 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 31737: contig of 31737 bp in length
31738 31837: gap of 100 bp
31838 36585: contig of 4748 bp in length
36586 36586: gap of 100 bp
36586 40932: contig of 4247 bp in length
40932 41032: gap of 100 bp
41032 44827: contig of 3795 bp in length
44827 44928: gap of 100 bp
44928 60753: contig of 15832 bp in length
60753 60760: gap of 100 bp
60760 73424: contig of 12565 bp in length
73424 73425: gap of 100 bp
73425 116324: contig of 42800 bp in length
116324 116424: gap of 100 bp
116424 142656: contig of 26232 bp in length
142656 142757: gap of 100 bp
142757 163746: contig of 20990 bp in length.

FEATURES

source

1. .163746
/organism="Mus musculus"
/mol_type="genomic DNA"
/db_xref="taxon:10090"
/clone="RP24-160E14"
/clone_lib="RPC1-24 Male Mouse BAC"
1. .31737
/note="assembly_fragment"
clone_end:SP6
vector_side:left
31838. .36585
/note="assembly_fragment"
36586. .40932
/note="assembly_fragment"
41033. .44827
/note="assembly_fragment"
44928. .60759
/note="assembly_fragment"

misc_feature

misc_feature

misc_feature

misc_feature

misc_feature

misc_feature

misc_feature

misc_feature

misc_feature

misc_feature

60860. .73424

/note="assembly_fragment"

73525. .116324

/note="assembly_fragment"

116425. .142656

/note="assembly_fragment"

142757. .163746

/note="assembly_fragment"

clone_end:T7

vector_side:right"

BASE COUNT 44170 a 38132 c 38033 g 42611 t 800 others

ORIGIN

Query Match 84.8%; Score 17.8; DB 2; Length 163746;

Best Local Similarity 90.5%; Pred. No. 4.7e+02;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGCCTGATGATGACGAGG 21

|||||

Db 159477 AGCTTGAATGATGACGAGG 159497

RESULT 9

AC109462/c

LOCUS

DEFINITION

AC109462 AC016225

AC109462.2 GI:19172840

KEYWORDS

HTG.

SOURCE

ORGANISM

Homo sapiens

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 174191)

DOE Joint Genome Institute.

AUTHORS

Sequencing of Human Chromosome 16

TITLE

Unpublished

REFERENCE

2 (bases 1 to 174191)

DOE Joint Genome Institute.

AUTHORS

Direct Submission

TITLE

Submitted (04-FEB-2002) Production Sequencing Facility, DOE Joint

Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA

3 (bases 1 to 174191)

DOE Joint Genome Institute.

AUTHORS

Direct Submission

TITLE

Submitted (06-MAR-2002) Production Sequencing Facility, DOE Joint

Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA

On or before Mar 6, 2002 this sequence version replaced

gi:10198460, gi:18483458.

COMMENT

Sequence Quality Assessment:

This entry has been annotated with sequence quality

estimates computed by the Phrap assembly program.

All manually edited bases have been reduced to quality zero.

Quality levels above 40 are expected to have less than

1 error in 10,000 bp.

Base-by-base quality values are not generally visible from the

GenBank flat file format but are available as part

of this entry's ASN.1 file.

Sequence Quality Assessment:

This entry has been annotated with sequence quality

estimates computed by the Phrap assembly program.

All manually edited bases have been reduced to quality zero.

Quality levels above 40 are expected to have less than

1 error in 10,000 bp.

Base-by-base quality values are not generally visible from the

GenBank flat file format but are available as part

of this entry's ASN.1 file.

FEATURES

source

Location/Qualifiers
1. .174191
/organism="Homo sapiens"
/mol_type="genomic DNA"

/db_xref="taxon:9606"

/chromosome="16"

/clone="RP11-26L20"

BASE COUNT 51311 a 36924 c 37255 g 48701 t

ORIGIN

Query Match 84.8%; Score 17.8; DB 9; Length 174191;

Best Local Similarity 90.5%; Pred. No. 4.7e+02;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAGCTGATTTCATCACCAGG 21

Db 39604 AAGCTGATTTCATCACCAGG 39584

RESULT 10

AC113908/c

LOCUS AC113908 177733 bp DNA linear HTG 15-NOV-2002

DEFINITION Rattus norvegicus clone CH230-396P4, WORKING DRAFT SEQUENCE.

AC113908

VERSION AC113908.9 GI:25008249

KEYWORDS HTG; HTGS PHASE2; HTGS DRAFT; HTGS_FULLTOP.

SOURCE Rattus norvegicus (Norway rat)

ORGANISM

Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

1 (bases 1 to 177733)

REFERENCE
Muzny,D.Marie., Metzker,M.Lee., Abramson,S., Adams,C., Alder,J.,
Allen,C., Allen,H., Alsbrooks,S., Amin,A., Anguiano,D.,
Anyalebechi,V., Aoyagi,A., Ayodeji,M., Baca,E., Baden,H.,
Baldwin,D., Bandaranaike,D., Barber,M., Barnstead,M., Benahmed,F.,
Biswal,K., Blair,J., Blankenburg,K., Blyth,P., Brown,M.,
Bryant,N., Buhay,C., Burch,P., Burrell,K., Calderon,E.,
Cardenas,V., Carter,K., Cavazos,I., Ceasar,H., Center,A.,
Chacko,J., Chavez,D., Chen,G., Chen,R., Chen,Y., Chen,Z., Chu,J.,
Cleveland,C., Cockrell,R., Cox,C., Coyie,M., Cree,A., D'Souza,L.,
Davila,M.L., Davis,C., Davy-Carroll,L., De Anda,C., Dederich,D.,
Delgado,O., Denison,S., Deramo,C., Ding,Y., Dinh,H., Divya,K.,
Draper,H., Dugan-Rocha,S., Dunn,A., Durbin,K., Duval,B., Eaves,K.,
Egan,A., Escotto,M., Eugene,C., Evans,C.A., Falls,T., Fan,G.,
Fernandez,S., Finley,M., Flagg,N., Forbes,L., Foster,M., Foster,P.,
Fraser,C.M., Gabisi,A., Ganta,R., Garcia,A., Garner,T., Garza,M.,
Gebregorgis,E., Geer,K., Gill,R., Grady,M., Guerra,W., Guevara,W.,
Gunaratne,P., Haaland,W., Hamli,C., Hamilton,C., Hamilton,K.,
Harvey,Y., Havlak,P., Hawes,A., Henderson,N., Hernandez,J.,
Hernandez,R., Hines,S., Hladun,S.L., Hodgson,A., Hoques,M.,
Hollins,B., Howells,S., Hulyk,S., Hume,J., Idlebird,D., Jackson,A.,
Jackson,L., Jacob,L., Jiang,H., Johnson,B., Johnson,R., Jolivet,A.,
Karpachy,S., Kelly,S., Kelly,S., Khan,Z., King,L., Kovar,C.,
Kowis,C., Kraft,C.L., Lebow,H., Levan,J., Lewis,L., Li,Z., Liu,J.,
Liu,J., Liu,W., Liu,Y., London,P., Longacre,S., Lopez,J.,
Lorensuhewa,L., Loulseghe,H., Lozada,R.J., Lu,X., Ma,J.,
Maheshwari,M., Mahindartine,M., Mahmoud,M., Malloy,K., Mangum,A.,
Mangum,B., Mapua,P., Martin,K., Martin,R., Martinez,E.,
Mawhinney,S., McLeod,M.P., McNeill,T.Z., Meenen,E.,
Milosavljevic,A., Miner,G., Minja,E., Montemayor,J., Moore,S.,
Morgan,M., Morris,K., Morris,S., Munidasa,M., Murphy,M., Nair,L.,
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Nwaokaleme,O., Okonou,G., Olampunsgoon,A., Pal,S., Parks,K.,
Pasternak,S., Paul,H., Perez,A., Perez,D., Pfamkoch,C.,
Plopper,F., Poinexter,A., Popovic,D., Primus,E., Pu,L.-L.,
Puzo,M., Quiroz,J., Rachlin,E., Reeves,K., Regier,M.A., Reigh,R.,
Reilly,B., Reilly,M., Ren,Y., Reuter,M., Richards,S., Riggs,F.,
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Sanders,W., Savary,G., Scherer,S., Scott,G., Shatsman,S., Shen,H.,
Shetty,J., Shvartsbeyn,A., Sisson,I., Sitter,C.D., Smajs,D.,
Sneed,A., Sodergren,E., Song,X.-Z., Sorelle,R., Sosa,J.,
Steimle,M., Strong,R., Sutton,A., Svatek,A., Tabor,P., Taylor,C.,
Taylor,T., Thomas,N., Thomas,S., Tingey,A., Trejos,Z., Usmani,K.,
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Williams,G., Willson,R., Wleczyk,R., Wooden,H., Worley,K.,

Wright,D., Wright,R., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V.,
Yu,F., Zhang,J., Zhou,J., Zhou,X., Zhao,S., Dunn,D., von
Niederhausern,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O.,
Weinstock,G. and Gibbs,R.A.

Direct Submission

Unpublished

2 (bases 1 to 177733)

Worley,K.C.

Direct Submission

Submitted (05-MAR-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA

3 (bases 1 to 177733)

Rat Genome Sequencing Consortium.

Direct Submission

Submitted (15-NOV-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA

On Nov 15, 2002 this sequence version replaced gi:23815563.

The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(<http://www.hgsc.bcm.tmc.edu/projects/rat/>). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.

----- Genome Center

Center: Baylor College of Medicine

Center code: BCM

Web site: <http://www.hgsc.bcm.tmc.edu/>

Contact: hgsc-help@bcm.tmc.edu

----- Project Information

Center project name: GTDH

Center clone name: CH230-396P4

----- Summary Statistics

Assembly program: Phrap; version 0.990329

Consensus quality: 165823 bases at least Q40

Consensus quality: 167723 bases at least Q30

Consensus quality: 168770 bases at least Q20

Estimated insert size: 172340; sum-of-contigs estimation

Quality coverage: 7x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).

* NOTE: This is a 'working draft' sequence. It currently

* consists of 1 contigs. Gaps between the contigs

* are represented as runs of N. The order of the pieces

* is believed to be correct as given, however the sizes

* of the gaps between them are based on estimates that have

* provided by the submitter.

* This sequence will be replaced

* by the finished sequence as soon as it is available and

* the accession number will be preserved.

* 177733: contig of 177733 bp in length.

FEATURES

source

1. 177733

/organism="Rattus norvegicus"

/mol_type="genomic DNA"

/db_xref="taxon:10116"

/clone="CH230-396P4"

342..1213

/note="clone_boundary"

clone_end:T7

site:

end sequence:B2175527"

complement(176718..177330)

/note="clone_boundary"

clone_end:Sp6

misc_feature

misc_feature


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site:
end sequence:BZ175528"
BASE COUNT 48653 a 37788 c 36881 g 46258 t 8153 others
ORIGIN

Query Match      84.8%; Score 17.8; DB 2; Length 177733;
Best Local Similarity 90.5%; Pred. No. 4.7e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAGCCTGATTGATGACGAGG 21
|||||
Db 72763 AAGCCTGATTGATGACGAG 72743
|||||

RESULT 11
AC128288/c
LOCUS AC128288 180353 bp DNA linear HTG 20-NOV-2002
DEFINITION Rattus norvegicus clone CH230-390F9, WORKING DRAFT SEQUENCE.
ACCESSION AC128288
VERSION AC128288.3 GI:25137732
KEYWORDS HTG; HTGS PHASE2; HTGS DRAFT; HTGS_FULLTOP.
SOURCE Rattus norvegicus (Norway rat)
ORGANISM Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
1 (bases 1 to 180353)
Muzny,D,Marie., Metzker,M, Lee., Abramson,S., Adams,C., Alder,J.,
Allen,C., Allen,H., Alsbrooks,S., Amin,A., Anguiano,D.,
Anyavebechi,V., Aoyagi,A., Ayodeji,M., Baca,E., Baden,H.,
Baldwin,D., Bandaranaike,D., Barber,M., Barnstead,M., Benahmed,F.,
Biswal,K., Blair,J., Blankenburg,K., Blyth,P., Brown,M.,
Bryant,N., Buhay,C., Burch,P., Burrell,K., Calderon,E.,
Cardenas,V., Carter,K., Cavazos,I., Cesar,H., Chen,A.,
Chacko,J., Chavez,D., Chen,G., Chen,R., Chen,Y., Chen,Z., Chu,J.,
Cleveland,C., Cockrell,R., Cox,C., Coyile,M., Cree,A., D'Souza,L.,
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Jackson,B., Jacob,L., Jiang,H., Johnson,B., Johnson,R., Jolivet,A.,
Karpachy,S., Kelly,S., Kelly,S., Khan,Z., King,L., Kovar,C.,
Kowis,C., Kraft,C.L., Lebow,H., Levan,J., Lewis,L., Li,Z., Liu,J.,
Liu,J., Liu,W., Liu,Y., London,P., Longacre,S., Lopez,J.,
Lorensuhewa,L., Loulseged,H., Lozado,R.J., Lu,X., Ma,J.,
Maheshwari,M., Mahindartne,M., Mahmoud,M., Malloy,K., Mangum,A.,
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Malosavljevic,A., Miner,G., Min'A.E., Montemayor,J., Moore,S.,
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Pasternak,S., Paul H.H., Perez,A., Perez,L., Pfannkoch,C.,
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Puazo,M., Quiroz,J., Rachlin,E., Reeves,K., Regier,M.A., Reigh,R.,
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Rives,C., Rodkey,T., Rojas,A., Rose,M., Rose,R., Ruiz,S.J.,
Sanders,W., Savary,G., Scherer,S., Scott,G., Shatsman,S., Shen,H.,
Shetty,J., Shvartsbeyn,A., Sisson,I., Sitter,C.D., Smajls,D.,
Sneed,A., Sodergren,E., Song,X.-Z., Sorelle,R., Sosa,J.,
Sreemle,M., Strong,R., Sutton,A., Svatek,A., Tabor,P., Taylor,C.,
Taylor,T., Thomas,N., Thomas,S., Tingey,A., Trejos,Z., Usmani,K.,
Valas,R., Vera,V., Villanasa,D., Waldron,L., Walker,B., Wang,J.,
Wang,Q., Wang,S., Warren,J., Warren,R., Wei,X., White,F.,
Williams,G., Willson,R., Wleczyk,R., Wooden,H., Worley,K.,
Wright,D., Wright,R., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V.,

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Yu,F., Zhang,J., Zhou,J., Zhou,X., Zhao,S., Dunn,D., von
Weinstock,G. and Gibbs,R.A.
Direct Submission
Unpublished
2 (bases 1 to 180353)
Worley,K.C.
Direct Submission
Submitted (19-JUL-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 180353)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (20-NOV-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GYDD
Center clone name: CH230-390F9
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 158606 bases at least Q40
Consensus quality: 170052 bases at least Q30
Consensus quality: 171200 bases at least Q20
Estimated insert size: 171038; sum-of-contigs estimation
Quality coverage: 8x in Q20 bases; sum-of-contigs estimation
-----
* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 1 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* provided by the submitter.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
* 1 180353: contig of 180353 bp in length.
* Location/Qualifiers
source
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/organism="Rattus norvegicus"
/mol_type="genomic DNA"
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misc_feature
1..1070
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2565..4033
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misc_feature
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BASE COUNT 48322 a 37960 c 37295 g 48481 t 8295 others
ORIGIN

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Query Match      84.8%; Score 17.8; DB 2; Length 180353;
Best Local Similarity 90.5%; Pred. No. 4.7e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAGCTGATTGATGACGAGG 21
Db 30172 AAGCTGATTGATGACGAG 30152

RESULT 12
AC117856/c
LOCUS AC117856
DEFINITION Rattus norvegicus clone CH230-327H18, WORKING DRAFT SEQUENCE, 3
unordered pieces.
ACCESSION AC117856
VERSION AC117856.4 GI:23814006
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.
SOURCE Rattus norvegicus (Norway rat)
ORGANISM Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
REFERENCE 1 (bases 1 to 186118)
AUTHORS Muzny,D., Maris., Metzker,M., Lee., Abranzon,S., Adams,C., Alder,J.,
Allen,C., Allen,H., Alsbrooks,S., Amin,A., Anguiano,D.,
Anyalebechi,V., Aoyagi,A., Ayodeji,M., Baca,E., Baden,H.,
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Chacko,J., Chavez,D., Chen,G., Chen,R., Chen,Y., Chen,Z., Chu,J.,
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Davila,M.L., Davis,C., Davy-Carroll,L., De Anda,C., Dederich,D.,
Delgado,O., Denison,S., Deramo,C., Ding,Y., Dinh,H., Divya,K.,
Draper,H., Dugan-Rocha,S., Dunn,A., Durbin,K., Duval,B., Eaves,K.,
Egan,A., Escotto,M., Eugene,C., Evans,C.A., Falls,T., Fan,G.,
Fernandez,S., Finley,M., Flagg,N., Forbes,L., Foster,M., Foster,P.,
Fraser,C.M., Gabisi,A., Ganta,R., Garcia,A., Garner,T., Garza,M.,
Gregorogis,E., Geer,K., Gill,R., Grady,M., Guerra,W., Guevara,W.,
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Liu,J., Liu,W., Liu,Y., London,P., Longacre,S., Lopez,J.,
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Mawhinney,S., McLeod,M.P., McNeill,T.Z., Meenen,E.,
Milosavljevic,A., Miner,G., Minja,E., Montemayor,J., Moore,S.,
Morgan,M., Morris,K., Morris,S., Munidasa,M., Murphy,M., Nair,L.,
Nankervis,C., Neal,D., Newton,N., Nguyen,N., Norris,S., Parks,K.,
Nwaekelameh,O., Okwono,G., Olarunpagoon,A., Pal,S., Parks,K.,
Pasternak,S., Paul,H., Perez,A., Perez,L., Pfankuch,C.,
Plopper,F., Poindexter,A., Popovic,D., Primus,E., Pu,L.-L.,
Puazo,M., Quiroz,J., Rachlin,E., Reeves,K., Regier,M.A., Reigh,R.,
Reilly,B., Reilly,M., Ren,Y., Reuter,M., Richards,S., Riggs,P.,
Rives,C., Rodkey,T., Rojas,A., Rose,M., Rose,R., Ruiz,S.,
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Yu,F., Zhang,J., Zhou,J., Zhou,X., Zhao,S., Zhao,S., Dunn,D., von
Niederhausern,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O.,
Weinstock,G. and Gibbs,R.A.
Direct Submission

```

JOURNAL

Unpublished

2 (bases 1 to 186118)

Worley,K.C.

Direct Submission

JOURNAL

Submitted (11-APR-2002) Human Genome Sequencing Center, Department

of Molecular and Human Genetics, Baylor College of Medicine, One

Baylor Plaza, Houston, TX 77030, USA

3 (bases 1 to 186118)

Rat Genome Sequencing Consortium.

Direct Submission

JOURNAL

Submitted (11-OCT-2002) Human Genome Sequencing Center, Department

of Molecular and Human Genetics, Baylor College of Medicine, One

Baylor Plaza, Houston, TX 77030, USA

On Oct 11, 2002 this sequence version replaced gi:21745720.

The sequence in this assembly is a combination of BAC based reads

and whole genome shotgun sequencing reads assembled using Atlas

(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described

in the feature table below represents a scaffold in the Atlas

assembly (a 'contig-scaffold'). Within each contig-scaffold,

individual sequence contigs are ordered and oriented, and separated

by sized gaps filled with Ns to the estimated size. The sequence

may extend beyond the ends of the clone and there may be sequence

contigs within a contig-scaffold that consist entirely of whole

genome shotgun sequence reads. Both end sequences and whole genome

shotgun sequence only contigs will be indicated in the feature

table.

----- Genome Center

Center: Baylor College of Medicine

Center code: BCM

Web site: http://www.hgsc.bcm.tmc.edu/

Contact: hgsc-help@bcm.tmc.edu

----- Project Information

Center project name: GTTF

Center clone name: CH230-327H18

----- Summary Statistics

Assembly program: Phrap; version 0.990329

Consensus quality: 169552 bases at least Q40

Consensus quality: 171519 bases at least Q30

Consensus quality: 172461 bases at least Q20

Estimated insert size: 173910; sum-of-contigs estimation

Quality coverage: 7x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length

* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).

* NOTE: This is a 'working draft' sequence. It currently

* consists of 3 contigs. The true order of the pieces

* is not known and their order in this sequence record is

* arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence

* as soon as it is available and the accession number will

* be preserved.

* 1 182244: contig of 182244 bp in length

* 182345 182344: gap of unknown length

* 182345 182442: contig of 1898 bp in length

* 184243 184342: gap of unknown length

* 184343 186118: contig of 1776 bp in length.

* Location/Qualifiers

1. 186118

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/mol_type="genomic DNA"

/db_xref="taxon:10116"

/clones="CH230-327H18"

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/note="wgs end extension

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complement(5228..6123)

/note="clone boundary

clone_end:T7"

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end_sequence:EXA0145TJ"

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/note="clone_boundary

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FEATURES

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misc_feature

misc_feature

misc_feature

TITLE

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BASE COUNT 49746 a 33747 c 35829 g 53877 t 12919 others
ORIGIN
Query Match 84.8%; Score 17.8; DB 2; Length 186118;
Best Local Similarity 90.5%; Pred. No. 4.7e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGCCCTGATGATGACACAGG 21
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Db 164995 AGCCCTGATGATGACACAGT 164975

RESULT 13
AC111040
LOCUS Mus musculus clone RP24-536C5, WORKING DRAFT SEQUENCE, 11 ordered
pieces.
AC111040 194454 bp DNA linear HTG 21-AUG-2002
HTG; HTGS PHAS2; HTGS DRAFT; HTGS_FULLTOP.
Mus musculus (house mouse)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
Birren,B., Nusbaum,C. and Lander,E.
Mus musculus, clone RP24-536C5
Unpublished
2 (bases 1 to 194454)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N.,
Anderson,S., Barna,N., Bastien,V., Boguslavsky,L., Boukhgalter,B.,
Brown,A., Camarata,J., Campopiano,A., Chang,J., Chazaro,B.,
Choepe,l., Colangelo,M., Collins,S., Collymore,A., Cook,A.,
Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S.,
Ferreira,P., FitzHugh,W., Gage,D., Galagan,J., Gardyna,S.,
Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N.,
Hagos,B., Horton,L., Hulme,W., Illiev,I., Johnson,R., Jones,C.,
Kamat,A., Karatas,A., Kells,C., LaRocque,K., Lamazares,R.,
Landers,T., Lehoczy,J., Levine,R., Liu,G., Maclean,C.,
Macdonald,P., Major,J., Marquis,N., Matthews,C., McCarthy,M.,
McSwan,P., McKernan,K., Meldrim,J., Meneus,L., Mihova,T.,
Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C.,
Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J.,
Peterson,K., Phunkhang,P., Pierre,N., Pollara,V., Raymond,C.,
Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P., Roman,J.,
Rosetti,W., Roy,A., Santos,R., Schauer,S., Schupback,R., Seaman,S.,
Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
Topham,K., Travers,M., Travis,N., Trigilio,J., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,
Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (18-FEB-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 194454)
Birren,B., Nusbaum,C., Lander,E., Allen,N., Anderson,S.,
Barna,N., Bastien,V., Bloom,T., Boguslavsky,L., Boukhgalter,B.,
Camarata,J., Chang,J., Chazaro,B., Choepe,l., Collymore,A.,
Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., FitzGerald,M., Gage,D., Galagan,J.,
Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B.,
Horton,L., Hulme,W., Illiev,I., Johnson,R., Jones,C., Kamat,A.,
Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K.,
Liu,G., Maclean,C., Macdonald,P., Major,J., Matthews,C.,
McCarthy,M., Meldrim,J., Meneus,L., Mihova,T., Mlenga,V.,
Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rieback,M.,
Roman,J., Rosetti,W., Roy,A., Santos,R., Schauer,S., Schupback,R.,
Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
Tesfaye,S., Theodore,J., Topham,K., Travers,M., Travis,N.,
Trigilio,J., Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X.,
Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (21-AUG-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Aug 21, 2002 this sequence version replaced gi:20806337.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L19423
Center clone name: 536_C5
----- Summary Statistics
Sequencing vector: Plasmid; n/a; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 191374 bases at least Q40
Consensus quality: 192313 bases at least Q30
Consensus quality: 192832 bases at least Q20
Insert size: 182000; agarose-fp
Insert size: 193454; sum-of-contigs
Quality coverage: 10.6 in Q20 bases; agarose-fp
Quality coverage: 10.0 in Q20 bases; sum-of-contigs
-----
* NOTE: This is a 'working draft' sequence. It currently
* consists of 11 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* been provided by the submitter.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
* 1 3686: contig of 3686 bp in length
* 3687 3786: gap of 100 bp
* 3787 11203: contig of 7417 bp in length
* 11204 11303: gap of 100 bp
* 11304 22134: contig of 10891 bp in length
* 22195 22294: gap of 100 bp
* 22295 32225: contig of 9931 bp in length
* 32226 32325: gap of 100 bp
* 32326 39995: contig of 7670 bp in length
* 39996 40095: gap of 100 bp
* 40096 57751: contig of 17656 bp in length
* 57752 57851: gap of 100 bp
* 57852 69355: contig of 11503 bp in length
* 69356 69454: gap of 100 bp
* 69455 87475: contig of 18021 bp in length
* 87476 87575: gap of 100 bp
* 87576 118208: contig of 30633 bp in length
* 118209 118308: gap of 100 bp
* 118309 158787: contig of 40479 bp in length
* 158788 158888: gap of 100 bp
* 158889 194454: contig of 35567 bp in length.
* Location/Qualifiers
* 1. 194454
* /organism="Mus musculus"
* /mol_type="genomic DNA"
* /db_xref="taxon:10090"
* /clone="RP24-536C5"
* /clone_lib="RPC1-24 Male Mouse BAC"
FEATURES
source

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misc_feature 1. 3686
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 /note="assembly_fragment"
 misc_feature 11304. 22194
 /note="assembly_fragment"
 misc_feature 22295. 32225
 /note="assembly_fragment"
 misc_feature 32326. 33995
 /note="assembly_fragment"
 misc_feature 40096. 57751
 /note="assembly_fragment"
 misc_feature 57852. 69354
 /note="assembly_fragment"
 misc_feature 69455. 87475
 /note="assembly_fragment"
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 misc_feature 158888. 194454
 /note="assembly_fragment"

BASE COUNT 53011 a 45075 c 45339 g 50027 t 1002 others
 ORIGIN

Query Match 84.8%; Score 17.8; DB 2; Length 194454;
 Best Local Similarity 90.5%; Pred. No. 4.7e+02;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAGCTGATTGATGACGAGG 21

Db 134728 AAGCTGATTGATGACGAGG 134748

RESULT 14
 AL691424/c
 LOCUS AL691424 194520 bp DNA linear ROD 26-JUL-2002
 DEFINITION Mouse DNA sequence from clone RP23-219K12 on chromosome X, complete
 sequence.
 ACCESSION AL691424.7 GI:22002699
 VERSION AL691424.7
 KEYWORDS HTG.
 SOURCE Mus musculus (house mouse)
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Euthera; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 194520)
 Wray.P.
 Direct Submission
 Submitted (26-JUL-2002) Wellcome Trust Sanger Institute, Hinxton,
 Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
 humquery@sanger.ac.uk
 On Jul 29, 2002 this sequence version replaced gi:21912679.
 COMMENT
 ----- Genome Center
 Center: Wellcome Trust Sanger Institute
 Center code: SC
 Web site: <http://www.sanger.ac.uk>
 Contact: humquery@sanger.ac.uk

During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above. This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. The following abbreviations are used to associate primary accession numbers given

in the feature table with their source databases: Em., EMBL; Sw., SWISSPROT; Tr., TREMBL; Wp., WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep RP23-219K12 is from the RPCI-23 Mouse PAC Library constructed by the group of Pieter de Jong. For further details see <http://www.chori.org/bacpac/home.htm>
 VECTOR: pBACe3.6.

FEATURES
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 1. 194520
 /organism="Mus musculus"
 /mol_type="genomic DNA"
 /db_xref="taxon:10090"
 /chromosome="X"
 /clone="RP23-219K12"
 /clone_lib="RPCI-23"
 BASE COUNT 55728 a 38763 c 37184 g 62845 t
 ORIGIN

Query Match 84.8%; Score 17.8; DB 10; Length 194520;
 Best Local Similarity 90.5%; Pred. No. 4.7e+02;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAGCTGATTGATGACGAGG 21

Db 42410 ATGCTGATTGATGACGAGG 42390

RESULT 15
 AC123058
 LOCUS AC123058 194636 bp DNA linear ROD 02-OCT-2002
 DEFINITION Mus musculus chromosome 19 clone RP23-188J8, complete sequence.
 ACCESSION AC123058
 VERSION AC123058.3 GI:23462991
 KEYWORDS HTG.
 SOURCE Mus musculus (house mouse)
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Euthera; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 194636)
 McPherson,J.D. and Waterston,R.H.
 The sequence of Mus musculus clone
 Unpublished
 REFERENCE 2 (bases 1 to 194636)
 McPherson,J.D. and Waterston,R.H.
 Direct Submission
 TITLE Direct Submission
 JOURNAL Submitted (27-MAY-2002) Genome Sequencing Center, 4444 Forest Park
 Parkway, St. Louis, MO 63108, USA
 3 (bases 1 to 194636)
 McPherson,J.D. and Waterston,R.H.
 Direct Submission
 TITLE Direct Submission
 JOURNAL Submitted (25-AUG-2002) Genome Sequencing Center, 4444 Forest Park
 Parkway, St. Louis, MO 63108, USA
 4 (bases 1 to 194636)
 McPherson,J.D. and Waterston,R.H.
 Direct Submission
 TITLE Direct Submission
 JOURNAL Submitted (02-OCT-2002) Genome Sequencing Center, 4444 Forest Park
 Parkway, St. Louis, MO 63108, USA
 On Oct 2, 2002 this sequence version replaced gi:22475873.

----- Genome Center -----
 Center: Washington University Genome Sequencing Center
 Center code: WUGSC
 Web site: <http://genome.wustl.edu/gsc/index.shtml>
 Contact: submissions@wustl.edu
 ----- Project Information -----
 Center project name: M_BA0188J08

FEATURES
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 1. 194636
 /organism="Mus musculus"
 /mol_type="genomic DNA"
 /db_xref="taxon:10090"

/chromosome="19"
/clone="RP23-188J8"
BASE COUNT 50027 a 45155 c 46196 g 53258 t
ORIGIN
Query Match 84.8%; Score 17.8; DB 10; Length 194636;
Best Local Similarity 90.5%; Pred. No. 4.7e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 AGCCTGATTGATGACCGGG 21
Db 50616 AGCCTGAATGATGACGAGGG 50636

Search completed: February 19, 2004, 22:22:05
Job time : 1601.5 secs


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Query Match
Best Local Similarity 70.5%; Score 14.8; DB 13; Length 20;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 ATCCTCACCTTGTCCTC 20
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Db 20 ATCCTCACCTTGTCCTC 3

RESULT 2
US-10-171-319-73
; Sequence 73, Application US/10171319
; Publication No. US20030157633A1
; GENERAL INFORMATION:
; APPLICANT: Ardem Patapoutian
; APPLICANT: Andrea Peier
; APPLICANT: Peter McIntyre
; APPLICANT: Stuart Bevan
; APPLICANT: Chuansheng Song
; APPLICANT: Pamposh Ganju
; TITLE OF INVENTION: VANILLOID RECEPTOR-RELATED NUCLEIC ACIDS
; TITLE OF INVENTION: AND POLYPEPTIDES
; FILE REFERENCE: 4-32048A
; CURRENT APPLICATION NUMBER: US/10/171,319
; CURRENT FILING DATE: 2002-10-24
; PRIOR APPLICATION NUMBER: 60/297,835
; PRIOR FILING DATE: 2001-06-13
; PRIOR APPLICATION NUMBER: 60/351,238
; PRIOR FILING DATE: 2002-01-22
; PRIOR APPLICATION NUMBER: 60/352,914
; PRIOR FILING DATE: 2002-01-29
; PRIOR APPLICATION NUMBER: 60/357,161
; PRIOR FILING DATE: 2002-02-12
; PRIOR APPLICATION NUMBER: 60/381,086
; PRIOR FILING DATE: 2002-05-15
; PRIOR APPLICATION NUMBER: 60/381,739
; PRIOR FILING DATE: 2002-05-16
; NUMBER OF SEQ ID NOS: 114
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 73
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide primer
US-10-171-319-73

Query Match
Best Local Similarity 70.5%; Score 14.8; DB 13; Length 20;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 ATCCTCACCTTGTCCTC 20
    |||||
Db 1 ATCCTCACCTTGTCCTC 18

RESULT 3
US-09-865-879-29
; Sequence 29, Application US/09865879
; Publication No. US20030180707A1
; GENERAL INFORMATION:
; APPLICANT: Roninbon, Igor
; APPLICANT: Dokmanovic, Milos
; APPLICANT: Chang, Bey-Dih
; TITLE OF INVENTION: REAGENTS AND METHODS FOR IDENTIFYING AND MODULATING EXPRESSION OF
; TITLE OF INVENTION: REGULATED BY RETINOIDS
; FILE REFERENCE: 99,216-H
; CURRENT APPLICATION NUMBER: US/09/865,879
; CURRENT FILING DATE: 2001-05-25
; PRIOR APPLICATION NUMBER: 60/207,535
; PRIOR FILING DATE: 2000-05-26
; NUMBER OF SEQ ID NOS: 44
; SOFTWARE: PatentIn version 3.0
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; SEQ ID NO 29
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc.feature
; OTHER INFORMATION: Antisense primer for P28 alpha
US-09-865-879-29

Query Match
Best Local Similarity 67.6%; Score 14.2; DB 13; Length 20;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TCATCTCACCTTGTCCT 19
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Db 2 TCATCTCACCTTGTCCT 20

RESULT 4
US-09-735-995-47/c
; Sequence 47, Application US/09735995
; Patent No. US20010034024A1
; GENERAL INFORMATION:
; APPLICANT: Keating, Mark T.
; APPLICANT: Splawski, Igor
; TITLE OF INVENTION: MUTATIONS IN AND GENOMIC STRUCTURE OF HERG - A LONG QT
; TITLE OF INVENTION: SYNDROME GENE
; FILE REFERENCE: 2323-136
; CURRENT APPLICATION NUMBER: US/09/735,995
; CURRENT FILING DATE: 2000-12-14
; PRIOR APPLICATION NUMBER: 09/226,012
; PRIOR FILING DATE: 1999-01-06
; NUMBER OF SEQ ID NOS: 116
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 47
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Homo sapiens
; OTHER INFORMATION:
US-09-735-995-47

Query Match
Best Local Similarity 66.7%; Score 14; DB 9; Length 20;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 CCTCACCTTGTCCT 18
    |||||
Db 16 CCTCACCTTGTCCT 3

RESULT 5
US-10-349-143-10433
; Sequence 10433, Application US/10349143
; Publication No. US20040005564A1
; GENERAL INFORMATION:
; APPLICANT: Cohen, Daniel
; APPLICANT: Blumenfeld, Marta
; APPLICANT: Chumakov, Ilya
; TITLE OF INVENTION: Biallelic markers for use in constructing a high density...
; FILE REFERENCE: GENSET.020CP1
; CURRENT APPLICATION NUMBER: US/10/349,143
; CURRENT FILING DATE: 2003-01-21
; PRIOR APPLICATION NUMBER: US/09/422,978
; PRIOR FILING DATE: 1999-10-20
; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: US 09/298,850
; PRIOR FILING DATE: EARLIER FILING DATE: 1999-04-21
; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: US 60/109,732
; PRIOR FILING DATE: EARLIER FILING DATE: 1998-11-23
; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: US 60/082,614
; PRIOR FILING DATE: EARLIER FILING DATE: 1998-04-21
; NUMBER OF SEQ ID NOS: 11796
; SEQ ID NO 10433
; LENGTH: 21
; TYPE: DNA
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; ORGANISM: Homo Sapiens
; FEATURE:
; NAME/KEY: primer_bind
; LOCATION: 1..21
; OTHER INFORMATION: downstream amplification primer 99-11851 for SEQ 2568, in complement
US-10-349-143-10433

Query Match      63.8%; Score 13.4; DB 12; Length 21;
Best Local Similarity 93.3%; Pred. No. 1.6e+04;
Matches 14; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 6 CTCACCCCTTGTCCTC 20
Db 7 CTCACCCCTTGTCCTC 21

RESULT 6
US-09-953-047-23
; Sequence 23, Application US/09953047
; Publication No. US20030087854A1
; GENERAL INFORMATION:
; APPLICANT: Brett P. Monia
; APPLICANT: Jacqueline Wyatt
; TITLE OF INVENTION: ANTISENSE MODULATION OF FIBROBLAST GROWTH FACTOR RECEPTOR 3 EXPRESSION
; FILE REFERENCE: RTS-0157
; CURRENT APPLICATION NUMBER: US/09/953,047
; CURRENT FILING DATE: 2001-09-10
; NUMBER OF SEQ ID NOS: 95
; SEQ ID NO 23
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Antisense Oligonucleotide
US-09-953-047-23

Query Match      62.9%; Score 13.2; DB 11; Length 20;
Best Local Similarity 83.3%; Pred. No. 2e+04;
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 4 TCCTCACCCCTTGTCCTCA 21
Db 2 TCCACACCTGTGTCTCTCA 19

RESULT 7
US-10-128-714-4014/c
; Sequence 4014, Application US/10128714
; Publication No. US20030119013A1
; GENERAL INFORMATION:
; APPLICANT: Jiang, Bo
; APPLICANT: Hu, Wenqi
; APPLICANT: Tishkoff, Daniel
; APPLICANT: Zamudio, Carlos
; APPLICANT: Broshkin, Alexey M
; APPLICANT: Lemieux, Sebastien M
; TITLE OF INVENTION: Identification of Essential Genes in Aspergillus fumigatus and
; TITLE OF INVENTION: Methods of Use
; FILE REFERENCE: 10182-018-999
; CURRENT APPLICATION NUMBER: US/10/128,714
; CURRENT FILING DATE: 2002-04-23
; PRIOR APPLICATION NUMBER: US 60/285,697
; PRIOR FILING DATE: 2001-04-23
; PRIOR APPLICATION NUMBER: US 60/287,066
; PRIOR FILING DATE: 2001-04-27
; PRIOR APPLICATION NUMBER: US 60/295,890
; PRIOR FILING DATE: 2001-06-05
; PRIOR APPLICATION NUMBER: US 60/303,899
; PRIOR FILING DATE: 2001-07-09
; PRIOR APPLICATION NUMBER: US 60/316,362
; PRIOR FILING DATE: 2001-08-31
; NUMBER OF SEQ ID NOS: 8603
; SOFTWARE: PatentIn version 3.1
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; SEQ ID NO 4014
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Aspergillus fumigatus
US-10-128-714-4014

Query Match      62.9%; Score 13.2; DB 15; Length 20;
Best Local Similarity 83.3%; Pred. No. 2e+04;
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 2 CATCCTCACCCCTTGTCCT 19
Db 19 CATACCCATCCTTGTCCT 2

RESULT 8
US-10-061-201-866/c
; Sequence 866, Application US/10061201
; Publication No. US20030166229A1
; GENERAL INFORMATION:
; APPLICANT: Shannon, Mark
; TITLE OF INVENTION: HUMAN POSH-LIKE PROTEIN 1
; FILE REFERENCE: PB0178
; CURRENT APPLICATION NUMBER: US/10/061,201
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00666
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00670
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/328,205
; PRIOR FILING DATE: 2001-10-10
; NUMBER OF SEQ ID NOS: 4162
; SOFTWARE: Asomica Sequence Listing Engine
; SEQ ID NO 866
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-061-201-866

Query Match      61.0%; Score 12.8; DB 13; Length 17;
Best Local Similarity 87.5%; Pred. No. 3e+04;
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 4 TCCTCACCCCTTGTCCT 19
Db 17 TTCACACTCTTGTCCT 2

RESULT 9
US-10-061-201-867/c
; Sequence 867, Application US/10061201
; Publication No. US20030166229A1
; GENERAL INFORMATION:
; APPLICANT: Shannon, Mark
; TITLE OF INVENTION: HUMAN POSH-LIKE PROTEIN 1
; FILE REFERENCE: PB0178
; CURRENT APPLICATION NUMBER: US/10/061,201
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00666
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RESULT 13
US-09-922-364A-35/c
; Sequence 35, Application US/09922364A
; Patent No. US20020155531A1
; GENERAL INFORMATION:
; APPLICANT: Adelman, John P.
; Maylie, James
; Bord, Chris T.
;

Silvia, Christopher P.
TITLE OF INVENTION: Small and Intermediate Conductance, Calcium-Activated Potassium Channels and Uses Thereof

NUMBER OF SEQUENCES: 48
CORRESPONDENCE ADDRESS:
ADDRESS: Townsend and Townsend and Crew LLP
STREET: Two Embarcadero Center, Eighth Floor
CITY: San Francisco
STATE: California
COUNTRY: USA
ZIP: 94111-3834

WIF. 54111-5634
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patent In Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/09/922,364A
 FILING DATE: 03-Aug-2001
 CLASSIFICATION: <Unknown>

CLASSIFICATION: UNCLASSIFIED

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 09/254,590

FILING DATE: 10-Mar-1999

APPLICATION NUMBER: US 60/026,451

FILING DATE: 11-SEP-1996

APPLICATION NUMBER: US 60/040,052

FILING DATE: 07-MAR-1997

APPLICATION NUMBER: US 60/045,233

FILING DATE: 17-APR-1997

APPLICATION NUMBER: WO PCT/US97/16033

FILING DATE: 10-SEP-1997

FILING DATE: 10-SEP-1997
ATTORNEY/AGENT INFORMATION:
NAME: Weber, Kenneth A.
REGISTRATION NUMBER: 31,677
REFERENCE/DOCKET NUMBER: 014210-000730US
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 576-0200
TELEFAX: (415) 576-0300

INFORMATION FOR SEQ ID NO: 35:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 21 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: DNA
 SEQUENCE DESCRIPTION: SEQ ID NO: 35:
 2-364A-35

Match 61.0%; Score 12.8; DB 10; Length 21;
Local Similarity 87.5%; Pred. No. 2.9e+04;
14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

5 CCTACCCCTTGTCCTC 20
|||||
19 CCTACCGCTTGGCCTC 4

4
1-590-35/c
ce 35, Application US/09254590
No. US20020165379A1
REAL INFORMATION:
APPLICANT: Adelman, John P.

SILVIA, CHRISTOPHER P.
TITLE OF INVENTION: Small and Intermediate Conductance,
Calcium-Activated Potassium Channels and Uses Thereof
NUMBER OF SEQUENCES: 48
CORRESPONDENCE ADDRESS:
ADDRESSEE: Townsend and Townsend and Crew LLP
STREET: Two Embarcadero Center, Eighth Floor

CITY: San Francisco
STATE: California
COUNTRY: USA
ZIP: 94111-3834
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/254.590
FILING DATE: 10-Mar-1999
CLASSIFICATION: <Unknown>

PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 60/026,451
 FILING DATE: 11-SEP-1996
 APPLICATION NUMBER: US 60/040,052
 FILING DATE: 07-MAR-1997
 APPLICATION NUMBER: US 60/045,233
 FILING DATE: 17-APR-1997
 APPLICATION NUMBER: WO PCT/US97/16033
 FILING DATE: 10-SEP-1997

FILING DATE: 10-25F-1237
ATTORNEY/AGENT INFORMATION:
NAME: Weber, Kenneth A.
REGISTRATION NUMBER: 31,677
REFERENCE/DOCKET NUMBER: 014210-000730US
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 576-0260

TELEPHONE: (415) 576-0300
TELEFAX: (415) 576-0300
INFORMATION FOR SEQ ID NO: 35:
SEQUENCE CHARACTERISTICS:
LENGTH: 21 base pairs
TYPE: nucleic acid
STRAINEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
SEQUENCE DESCRIPTION: SEQ ID NO: 35:
1-590-35

Match	61.0%;	Score 12.8;	DB 10;	Length 21;
Local Similarity	87.5%;	Pred. No. 2.9e+04;		
14: Conservative	0;	Mismatches 2;	Indels 0;	Gaps 0;

5 CCTCACCCCTTGTCCTC 20
|||||
19 CCTCACGGTTGGCCTC 4

5
100-143-9636/c
Application 9636, Application US/10349143
Application No. US20040005584A1
INFORMATION:
CANT: Cohen, Daniel
CANT: Blumenfeld, Marta
CANT: Chumakov, Ilya
OF INVENTION: Biallelic markers for use in constructing a high density...
REFERENCE: GENSET.020CP1
FIRST APPLICATION NUMBER: US/10/349,143
FIRST FILING DATE: 2003-01-21
APPLICATION NUMBER: US/09/422,978

FILING DATE: 1999-10-20
 APPLICATION NUMBER: EARLIER APPLICATION NUMBER: US 09/298,850
 FILING DATE: EARLIER FILING DATE: 1999-04-21
 APPLICATION NUMBER: EARLIER APPLICATION NUMBER: US 60/109,732
 FILING DATE: EARLIER FILING DATE: 1998-11-23
 APPLICATION NUMBER: EARLIER APPLICATION NUMBER: US 60/082,614
 FILING DATE: EARLIER FILING DATE: 1998-04-21
 R OF SEQ ID NOS: 11796
 NO 9635
 ETH: 21
 : DNA
 NISM: Homo Sapiens

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; FEATURE:
; NAME/KEY: primer_bind
; LOCATION: 1..21
; OTHER INFORMATION: downstream amplification primer 99-6261 for SEQ 1771, in compleme
US-10-349-143-9636
Query Match      61.0%; Score 12.8; DB 12; Length 21;
Best Local Similarity 87.5%; Pred. No. 2.9e+04;
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY      5 CCTCACCCCTTGTCTC 20
Db      21 CCTCACCCCTTCTC 6
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Search completed: February 20, 2004, 04:09:30
Job time : 198 secs